

|    | Type | Hits | Search Text                                  |
|----|------|------|--|
| 1  | IS&R | 1    | ("6159495").PN.                              |
| 2  | IS&R | 1    | ("6752834").PN.                              |
| 3  | IS&R | 0    | ("2002013408").PN.                           |
| 4  | IS&R | 0    | ("2001018614").PN.                           |
| 5  | IS&R | 1    | ("5972385").PN.                              |
| 6  | IS&R | 0    | ("200183858").PN.                            |
| 7  | IS&R | 0    | ("2005124797").PN.                           |
| 8  | IS&R | 0    | ("2005124797").PN.                           |
| 9  | BRS  | 1    | (sulfonated same keratin) and hydroxyapatite |
| 10 | BRS  | 1    | (sulfonated with keratin) and hydroxyapatite |
| 11 | BRS  | 1    | (sulfonated near keratin) and hydroxyapatite |
| 12 | BRS  | 205  | sulfonated and keratin and hydroxyapatite    |
| 13 | BRS  | 0    | (sulfonated and keratin) near hydroxyapatite |
| 14 | IS&R | 1    | ("7148327").PN.                              |
| 15 | BRS  | 23   | keratin with hydroxyapatite                  |
| 16 | BRS  | 919  | keratin and hydroxyapatite                   |
| 17 | BRS  | 1    | (sulfonated and keratin) same hydroxyapatite |
| 18 | BRS  | 1    | keratin near hydroxyapatite                  |
| 19 | BRS  | 141  | keratin same hydroxyapatite                  |
| 20 | BRS  | 1    | (sulfonated and keratin) with hydroxyapatite |
| 21 | IS&R | 0    | ("2002177903").PN.                           |
| 22 | IS&R | 0    | ("2002177903A1").PN:                         |

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| Set                   | Items | Description          |
|-----------------------|-------|----------------------|
| ? s keratin           | S1    | 75091 KERATIN        |
| ? s hydroxyapatite    | S2    | 74761 HYDROXYAPATITE |
| ? s s1 and s2         | 75091 | S1                   |
|                       | 74761 | S2                   |
|                       | S3    | 40 S1 AND S2         |
| ? s s3 and sulfonated | 40    | S3                   |
|                       | 15491 | SULFONATED           |
| ? t s3/9,k/1-10       | S4    | 0 S3 AND SULFONATED  |

3/9,K/1 (Item 1 from file: 5)  
DIALOG(R)File 5:Biosis Previews(R)  
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0016268770 BIOSIS NO.: 200600614165  
Structure of white rhinoceros (*Ceratotherium simum*) horn investigated by X-ray histology with implications computed tomography and for growth and external form  
AUTHOR: Hieronymus Tobin L (Reprint); Witmer Lawrence M; Ridgely Ryan C  
AUTHOR ADDRESS: Ohio Univ, Dept Sci Biol, Irvine Hall, Athens, OH 45701 USA  
\*\*USA  
AUTHOR E-MAIL ADDRESS: Th108702@ohiou.edu  
JOURNAL: Journal of Morphology 267 (10): p1172-1176 OCT 2006 2006  
ISSN: 0362-2525  
DOCUMENT TYPE: Article  
RECORD TYPE: Abstract  
LANGUAGE: English

ABSTRACT: The nasal and frontal horns of two individuals of *Ceratotherium simum* were examined by x-ray computed tomography (CT scanning), gross observation of sectioned horn, and light microscopy of histological sections of the horn tissue. CT scans of both sets of horns reveal a periodic banding pattern that is evident upon gross observation of

sections as darker bands of tissue. The overlap of these bands in both histological and CT slices suggests the presence of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely hydroxyapatite or octocalcium phosphate). The distribution of these two components in the horns is hypothesized to contribute to the differential wear patterns that produce the characteristic sweeping conical shape of rhinoceros horn from what otherwise (in the absence of wear and UV exposure) would be cylindrical blocks of constantly growing cornified papillary epidermis. Although extant rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of keratin -and-bone horns such as those of bovid artiodactyls, the tissue structures that make up the horn are strikingly convergent with other examples of papillary cornified epidermis found in horses, artiodactyls, cetaceans, and birds.

REGISTRY NUMBERS: 169799-44-4: keratin ; 10103-46-5: calcium phosphate  
DESCRIPTORS:

MAJOR CONCEPTS: Methods and Techniques; Integumentary System--Chemical Coordination and Homeostasis

BIOSYSTEMATIC NAMES: Rhinocerotidae--Perissodactyla, Mammalia, Vertebrata, Chordata, Animalia

ORGANISMS: Ceratotherium simum {white rhinoceros} (Rhinocerotidae)

ORGANISMS: PARTS ETC: frontal horn; nasal horn; papillary epidermis-- integumentary system

COMMON TAXONOMIC TERMS: Animals; Chordates; Mammals; Nonhuman Vertebrates ; Nonhuman Mammals; Perissodactyls; Vertebrates

CHEMICALS & BIOCHEMICALS: keratin ; melanin; calcium phosphate

METHODS & EQUIPMENT: light microscopy--laboratory techniques, imaging and microscopy techniques; X-ray computed tomography--laboratory techniques, imaging and microscopy techniques

CONCEPT CODES:

10064 Biochemistry studies - Proteins, peptides and amino acids

10069 Biochemistry studies - Minerals

18504 Integumentary system - Physiology and biochemistry

BIOSYSTEMATIC CODES:

86150 Rhinocerotidae

...ABSTRACT: of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely hydroxyapatite or octocalcium phosphate). The distribution of these two components in the horns is hypothesized to...

...rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of keratin -and-bone horns such as those of bovid artiodactyls, the tissue structures that make up...

...REGISTRY NUMBERS: keratin ;

DESCRIPTORS:

CHEMICALS & BIOCHEMICALS: keratin ;

3/9,K/2 (Item 2 from file: 5)

DIALOG(R) File 5:Biosis Previews(R)

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0015091467 BIOSIS NO.: 200400472696

Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation

AUTHOR: Tachibana Akira; Kaneko Sumika; Tanabe Toshizumi; Yamauchi Kiyoshi  
(Reprint)  
AUTHOR ADDRESS: Grad Sch EngnDept Appl and Bioappl ChemSumiyoshi Ku, Osaka  
City Univ, Sugimoto 3-3-138, Osaka, 5588585, Japan\*\*Japan  
AUTHOR E-MAIL ADDRESS: tatibana@bioa.eng.osaka-cu.ac.jp;  
Yamauchi@bioa.eng.osaka-cu.ac.jp  
JOURNAL: Biomaterials 26 (3): p297-302, 285 January 2005 2005  
MEDIUM: print  
ISSN: 0142-9612  
DOCUMENT TYPE: Article  
RECORD TYPE: Abstract  
LANGUAGE: English

**ABSTRACT:** Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium phosphate hybrid biomaterials were described. Firstly, the CaP-precipitated sponges were obtained by only the immersion of the carboxyl-sponges, chemically introduced with high amount of carboxyl groups on the sponges, in calcium and phosphate ions containing buffers such as PBS(+) for only 1-3 days. Neither sponge, introduced with amino or amido groups or non-treated, gave significant calcium phosphate precipitation. The carboxyl-sponges were mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast differentiation marker, suggested that both hybrid sponges, CaP-precipitated and trapped sponges, alter the differentiation pattern of preosteoblasts, MC3T3-E1. Copyright 2004 Elsevier Ltd. All rights reserved.

REGISTRY NUMBERS: 9001-78-9: alkaline phosphatase; 14127-61-8: calcium ion; 7758-87-4Q: calcium phosphate; 10103-46-5Q: calcium phosphate; 56271-99-9: gamma-carboxyglutamic acid; 1306-06-5: hydroxyapatite ; 169799-44-4: keratin ; 14265-44-2: phosphate ion

ENZYME COMMISSION NUMBER: EC 3.1.3.1: alkaline phosphatase

DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Cell Biology; Methods and Techniques; Skeletal System--Movement and Support

BIOSYSTEMATIC NAMES: Muridae--Rodentia, Mammalia, Vertebrata, Chordata, Animalia

ORGANISMS: MC3T3E1 cell line (Muridae)--mouse osteoblast cells

ORGANISMS: PARTS ETC: osteoblast--skeletal system, cultivation, differentiation

COMMON TAXONOMIC TERMS: Animals; Chordates; Mammals; Nonhuman Vertebrates ; Nonhuman Mammals; Rodents; Vertebrates

CHEMICALS & BIOCHEMICALS: PBS buffer; alkaline phosphatase--expression; calcium ion; calcium phosphate; carboxyl group; gamma-carboxyglutamic acid; hydroxyapatite ; keratin ; phosphate ion; wool keratin

METHODS & EQUIPMENT: cell fabrication--culturing techniques, laboratory techniques

MISCELLANEOUS TERMS: high density cell cultivation; keratin - hydroxyapatite hybrid sponge; osteoblast calcification

CONCEPT CODES:

02502 Cytology - General  
02506 Cytology - Animal  
10060 Biochemistry studies - General  
10064 Biochemistry studies - Proteins, peptides and amino acids  
10069 Biochemistry studies - Minerals  
10802 Enzymes - General and comparative studies: coenzymes  
18004 Bones, joints, fasciae, connective and adipose tissue - Physiology  
and biochemistry

BIOSYSTEMATIC CODES:

86375 Muridae

Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation

ABSTRACT: Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium...

...mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast...

...REGISTRY NUMBERS: hydroxyapatite ; ...

... keratin ;

DESCRIPTORS:

CHEMICALS & BIOCHEMICALS: ... hydroxyapatite ; ...

... keratin ; ...

...wool keratin

MISCELLANEOUS TERMS: ... keratin - hydroxyapatite hybrid sponge

3/9,K/3 (Item 3 from file: 5)

DIALOG(R) File 5:Biosis Previews(R)

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0013550978 BIOSIS NO.: 200200144489

Isolation of Thermoanaerobacter keratinophilus sp. nov., a novel thermophilic, anaerobic bacterium with keratinolytic activity

AUTHOR: Riessen Sabine (Reprint); Antranikian Garabed

AUTHOR ADDRESS: Institute of Technical Microbiology, Technical University Hamburg-Harburg, Kasernenstr. 12, D-21073, Hamburg, Germany\*\*Germany

JOURNAL: Extremophiles 5 (6): p399-408 December, 2001 2001

MEDIUM: print

ISSN: 1431-0651

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: Several thermophilic anaerobic bacteria with keratinolytic activity growing at temperatures between 50degreeC and 90degreeC were

isolated from samples collected on the island of Sao Miguel in the Azores (Portugal). On the basis of morphological, physiological, and 16S rDNA studies, the isolate 2KXI was identified as a new species of the genus *Thermoanaerobacter*, designated *Thermoanaerobacter keratinophilus*. This strain, which grows optimally at 70degreeC, pH 7.0, and 0.5% NaCl, is the first member of the genus *Thermoanaerobacter* that has been described for its ability to degrade native **keratin**. Around 70% of native wool was solubilized after 10 days of incubation under anaerobic conditions. The strain was shown to possess intracellular and extracellular proteases optimally active at 60degreeC, pH 7.0, and 85degreeC, pH 8.0, respectively. Keratin hydrolysis was demonstrated in vitro using a sodium dodecyl sulfate gel containing feather meal. The extracellular protease responsible for breaking down **keratin** fibers was purified to homogeneity in only one step by applying **hydroxyapatite** column chromatography. The enzyme belongs to the serine-type proteases and has a molecular mass of 135 kDa.

REGISTRY NUMBERS: 169799-44-4: **keratin** ; 9001-92-7: **protease**; 37259-58-8: **serine-type proteases**

DESCRIPTORS:

MAJOR CONCEPTS: Bacteriology; Methods and Techniques; Systematics and Taxonomy

BIOSYSTEMATIC NAMES: Irregular Nonsporing Gram-Positive Rods--Actinomycetes and Related Organisms, Eubacteria, Bacteria, Microorganisms

ORGANISMS: *Thermoanaerobacter* (Irregular Nonsporing Gram-Positive Rods); *Thermoanaerobacter keratinophilus* (Irregular Nonsporing Gram-Positive Rods)--isolation, new species, thermophilic anaerobic bacterium

COMMON TAXONOMIC TERMS: Bacteria; Eubacteria; Microorganisms

CHEMICALS & BIOCHEMICALS: **keratin**--hydrolysis; **protease**--extracellular, intracellular; **serine-type proteases**

METHODS & EQUIPMENT: 16S rDNA study {16S ribosomal DNA study}--analytical method, molecular genetic method; SDS-polyacrylamide gel electrophoresis {SDS-PAGE}--detection method; **hydroxyapatite** column chromatography--purification method

GEOGRAPHICAL NAME: Sao Miguel (Portugal, Europe) (Palearctic region)

MISCELLANEOUS TERMS: **keratinolytic activity**

CONCEPT CODES:

00504 General biology - Taxonomy, nomenclature and terminology

10064 Biochemistry studies - Proteins, peptides and amino acids

10802 Enzymes - General and comparative studies: coenzymes

30000 Bacteriology, general and systematic

31000 Physiology and biochemistry of bacteria

BIOSYSTEMATIC CODES:

08890 Irregular Nonsporing Gram-Positive Rods

...ABSTRACT: member of the genus *Thermoanaerobacter* that has been described for its ability to degrade native **keratin**. Around 70% of native wool was solubilized after 10 days of incubation under anaerobic conditions...  
...extracellular proteases optimally active at 60degreeC, pH 7.0, and 85degreeC, pH 8.0, respectively. Keratin hydrolysis was demonstrated in vitro using a sodium dodecyl sulfate gel containing feather meal. The extracellular protease responsible for breaking down **keratin** fibers was purified to homogeneity in only one step by applying **hydroxyapatite** column chromatography. The enzyme belongs to the serine-type proteases and has a molecular mass...  
...REGISTRY NUMBERS: **keratin** ;

DESCRIPTORS:

CHEMICALS & BIOCHEMICALS: keratin --

...METHODS & EQUIPMENT: hydroxyapatite column chromatography

3/9, K/4 (Item 4 from file: 5)

DIALOG(R) File 5:Biosis Previews(R)

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0010324311 BIOSIS NO.: 199698792144

The mineralization of crystalline inorganic components in Japanese serow horn

AUTHOR: Hashiguchi Kunio; Hashimoto Kenji

AUTHOR ADDRESS: Dep. Oral Surgery, Hamamatsu Univ. Sch. Med., 3600

Handa-cho, Hamamatsu, Shizuoka 431-31, Japan\*\*Japan

JOURNAL: Okajimas Folia Anatomica Japonica 72 (5): p235-244 1995 1995

ISSN: 0030-154X

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: English

ABSTRACT: The Japanese serow (*Capricornis crispus*) is protected as a special natural monument in Japan. The ring count of the soft X-ray photographs of Japanese serow horn was found to be a useful criteria to determine the ages exactly. The mineralization process in Japanese serow horn was examined microscopic, ICP and X-ray diffraction methods. The incremental lines appeared as light and dark layers in the section stained for fuchsin and methylen blue. Mineral depositions were observed among the keratin fibers, no matrix vesicle in the electron dense regions. X-ray diffraction pattern of crystalline inorganic components in Japanese serow horn was determined as beta-tricalcium phosphate (TCP), hydroxyapatite (HA) and unknown phase. ICP measurement was also carried out. The horn contained trace elements of K besides Na, Ca, Fe and P. The Ca/P molar was found to be 2.9. The ratio was much higher than the theoretical value of HA. Presumably, keratin was the seed which might be related to mineralization and higher Ca activity was detected in the initial phase of epitaxial growth. Analytical results of the measurement of trace elements in Japanese serow horn by using ICP method seemed to be correlated with the evaluation of environmental conditions. The present study indicated that the mineralization of Japanese serow horn directly related with deposition Ca-deficient HA among the keratin fibers.

REGISTRY NUMBERS: 7758-87-4: BETA-TRICALCIUM PHOSPHATE; 1306-06-5:

HYDROXYAPATITE ; 7440-09-7: POTASSIUM; 7440-23-5: SODIUM; 7440-70-2:

CALCIUM; 7439-89-6: IRON; 7723-14-0: PHOSPHORUS

DESCRIPTORS:

MAJOR CONCEPTS: Development; Integumentary System--Chemical Coordination and Homeostasis; Metabolism; Morphology; Radiology--Medical Sciences; Skeletal System--Movement and Support

BIOSYSTEMATIC NAMES: Bovidae--Artiodactyla, Mammalia, Vertebrata, Chordata, Animalia

ORGANISMS: *Capricornis crispus* (Bovidae)

COMMON TAXONOMIC TERMS: Animals; Artiodactyls; Chordates; Mammals; Nonhuman Vertebrates; Nonhuman Mammals; Vertebrates

CHEMICALS & BIOCHEMICALS: BETA-TRICALCIUM PHOSPHATE; HYDROXYAPATITE ; POTASSIUM; SODIUM; CALCIUM; IRON; PHOSPHORUS

MISCELLANEOUS TERMS: AGE; ANATOMY; BETA-TRICALCIUM PHOSPHATE; CALCIUM;

EPIDERMAL MINERALIZATION; EPITAXIAL GROWTH; HYDROXYAPATITE ; IRON;  
KERATIN FIBER; MICROSCOPY; MINERAL DEPOSITION; PHOSPHORUS; POTASSIUM;  
SODIUM; TRACE ELEMENT; X-RAY DIFFRACTION TECHNIQUE

CONCEPT CODES:

- 06504 Radiation biology - Radiation and isotope techniques
- 10060 Biochemistry studies - General
- 10064 Biochemistry studies - Proteins, peptides and amino acids
- 10069 Biochemistry studies - Minerals
- 11106 Anatomy and Histology - Radiologic anatomy
- 11108 Anatomy and Histology - Microscopic and ultramicroscopic anatomy
- 13002 Metabolism - General metabolism and metabolic pathways
- 13010 Metabolism - Minerals
- 13012 Metabolism - Proteins, peptides and amino acids
- 18002 Bones, joints, fasciae, connective and adipose tissue - Anatomy
- 18004 Bones, joints, fasciae, connective and adipose tissue - Physiology  
and biochemistry
- 18502 Integumentary system - Anatomy
- 18504 Integumentary system - Physiology and biochemistry
- 25508 Development and Embryology - Morphogenesis

BIOSYSTEMATIC CODES:

- 85715 Bovidae

...ABSTRACT: in the section stained for fuchsin and methylen blue. Mineral depositions were observed among the keratin fibers, no matrix vesicle in the electron dense regions. X-ray diffraction pattern of crystalline inorganic components in Japanese serow horn was determined as beta-tricalcium phosphate (TCP), hydroxyapatite (HA) and unknown phase. ICP measurement was also carried out. The horn contained trace elements

...  
...be 2.9. The ratio was much higher than the theoretical value of HA. Presumably, keratin was the seed which might be related to mineralization and higher Ca activity was detected...

...the mineralization of Japanese serow horn directly related with deposition Ca-deficient HA among the keratin fibers.

...REGISTRY NUMBERS: HYDROXYAPATITE ;

DESCRIPTORS:

CHEMICALS & BIOCHEMICALS: ... HYDROXYAPATITE ;

MISCELLANEOUS TERMS: ... HYDROXYAPATITE ; ...

... KERATIN FIBER

3/9,K/5 (Item 5 from file: 5)

DIALOG(R) File 5:Biosis Previews(R)

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0007376034 BIOSIS NO.: 199140018925

ESTABLISHMENT OF A HUMAN GINGIVAL EPITHELIAL CELL LINE WITH ACTIVITY OF KERATIN SYNTHESIS

AUTHOR: ISHIKAWA K (Reprint); MOCHII M; AGATA K; EGUCHI G

AUTHOR ADDRESS: DEP DEV BIOL, NATL INST BASIC BIOL, OKAZAKI 444, JPN\*\*JAPAN

JOURNAL: Development Growth and Differentiation 32 (4): p414 1990

CONFERENCE/MEETING: 23RD ANNUAL MEETING OF THE JAPANESE SOCIETY OF DEVELOPMENTAL BIOLOGISTS, HIROSHIMA, JAPAN, MAY 24-26, 1990. DEV GROWTH DIFFER.

ISSN: 0012-1592

DOCUMENT TYPE: Meeting

RECORD TYPE: Citation

LANGUAGE: ENGLISH

DESCRIPTORS: ABSTRACT HUMAN ORAL SURGERY IN-VITRO EXPERIMENTAL SYSTEMS  
HYDROXYAPATITE TISSUE CULTURE CLONAL SELECTION TECHNIQUES DEVELOPMENTAL  
RESEARCH ADHESION SPREADING GROWTH

DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Cell Biology;  
Dental and Oral System--Ingestion and Assimilation; Development;  
Methods and Techniques  
BIOSYSTEMATIC NAMES: Hominidae--Primates, Mammalia, Vertebrata, Chordata,  
Animalia  
COMMON TAXONOMIC TERMS: Animals; Chordates; Humans; Mammals; Primates;  
Vertebrates

CONCEPT CODES:

00520 General biology - Symposia, transactions and proceedings  
02508 Cytology - Human  
10054 Biochemistry methods - Proteins, peptides and amino acids  
10060 Biochemistry studies - General  
10064 Biochemistry studies - Proteins, peptides and amino acids  
11105 Anatomy and Histology - Surgery  
19001 Dental - General and methods  
19002 Dental - Anatomy  
19004 Dental - Physiology and biochemistry  
25504 Development and Embryology - Experimental  
25508 Development and Embryology - Morphogenesis  
32500 Tissue culture, apparatus, methods and media

BIOSYSTEMATIC CODES:

86215 Hominidae

**ESTABLISHMENT OF A HUMAN GINGIVAL EPITHELIAL CELL LINE WITH ACTIVITY OF  
KERATIN SYNTHESIS**

DESCRIPTORS: ABSTRACT HUMAN ORAL SURGERY IN-VITRO EXPERIMENTAL SYSTEMS  
HYDROXYAPATITE TISSUE CULTURE CLONAL SELECTION TECHNIQUES DEVELOPMENTAL  
RESEARCH ADHESION SPREADING GROWTH

3/9,K/6 (Item 6 from file: 5)

DIALOG(R) File 5:Biosis Previews(R)

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0004312443 BIOSIS NO.: 198478047850

**ASPECTS OF THE STRUCTURE AND COMPOSITION OF BALEEN AND SOME EFFECTS OF  
EXPOSURE TO PETROLEUM HYDRO CARBONS**

AUTHOR: ST AUBIN D J (Reprint); STINSON R H; GERACI J R

AUTHOR ADDRESS: WILDLIFE SECT, DEP PATHOL, ONTARIO VET COLL, UNIV GUELPH,  
GUELPH, ONT, CANADA N1G 2W1\*\*CANADA

JOURNAL: Canadian Journal of Zoology 62 (2): p193-198 1984

ISSN: 0008-4301

DOCUMENT TYPE: Article

RECORD TYPE: Abstract

LANGUAGE: ENGLISH

ABSTRACT: The structure and composition of baleen from 7 spp. of whales was  
studied using tensiometry, X-ray diffraction and elemental analysis.  
Baleen was composed principally of amorphous and .alpha.- keratin .

**Hydroxyapatite** (bone mineral, Ca<sub>10</sub> (PO<sub>4</sub>)<sub>6</sub> OH<sub>2</sub>) was present in all species. Certain elements, Mn, Cu, B, Fe and Ca were more highly concentrated in the fibers than in the matrix of the plate. The breaking strength of baleen plates from fin (*Balaenoptera physalus*), sei (*B. borealis*) and gray (*Eschrichtius robustus*) whales was comparable to that of buffalo horn, in the range of 2-9 times. 106 N .cntdot. m-2. The stiffness of baleen was somewhat less than that of other keratinized tissues. Treatment with 10% (vol/vol) trichloroacetic acid for 8 days removed most of the Ca salts, denatured .alpha.- keratin , and made fin whale plates stronger and stiffer. Exposure to gasoline for 1.5 h or 14 days, crude oil for 8 days, or tar for 21 days resulted in loss of trace elements from baleen and inconsistent changes in keratin organization. After tar exposure, fin whale baleen plates were stiffer and stronger. At sea, baleen would be relatively resistant to damage by spilled oil.

REGISTRY NUMBERS: 1306-06-5: HYDROXYLAPATITE; 7440-70-2: CALCIUM; 7439-89-6 : IRON; 7440-42-8: BORON; 7440-50-8: COPPER; 7439-96-5: MANGANESE

DESCRIPTORS: BALAENOPTERA-PHYSALUS BALAENOPTERA-BOREALIS  
ESCHRICHTIUS-ROBUSTUS KERATIN HYDROXYLAPATITE CALCIUM IRON BORON COPPER  
MANGANESE BREAKING STRENGTH/

DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Digestive System-- Ingestion and Assimilation; Pollution Assessment Control and Management ; Toxicology

BIOSYSTEMATIC NAMES: *Balaenopteridae*--Cetacea, Mammalia, Vertebrata, Chordata, Animalia; *Desmodontidae*--Chiroptera, Mammalia, Vertebrata, Chordata, Animalia

COMMON TAXONOMIC TERMS: Cetaceans; Animals; Bats; Chordates; Mammals; Nonhuman Vertebrates; Nonhuman Mammals; Vertebrates

CHEMICALS & BIOCHEMICALS: HYDROXYLAPATITE; CALCIUM; IRON; BORON; COPPER ; MANGANESE

CONCEPT CODES:

07512 Ecology: environmental biology - Oceanography  
07517 Ecology: environmental biology - Water research and fishery biology

10064 Biochemistry studies - Proteins, peptides and amino acids

10069 Biochemistry studies - Minerals

14002 Digestive system - Anatomy

22506 Toxicology - Environment and industry

37015 Public health - Air, water and soil pollution

BIOSYSTEMATIC CODES:

85810 *Balaenopteridae*

85850 *Desmodontidae*

...ABSTRACT: tensiometry, X-ray diffraction and elemental analysis. Baleen was composed principally of amorphous and .alpha.- keratin .

**Hydroxyapatite** (bone mineral, Ca<sub>10</sub> (PO<sub>4</sub>)<sub>6</sub> OH<sub>2</sub>) was present in all species. Certain elements, Mn, Cu...

...vol/vol) trichloroacetic acid for 8 days removed most of the Ca salts, denatured .alpha.- keratin , and made fin whale plates stronger and stiffer. Exposure to gasoline for 1.5 h...

...for 21 days resulted in loss of trace elements from baleen and inconsistent changes in keratin organization. After tar exposure, fin whale baleen plates were stiffer and stronger. At sea, baleen...

DESCRIPTORS: BALAENOPTERA-PHYSALUS BALAENOPTERA-BOREALIS  
ESCHRICHTIUS-ROBUSTUS KERATIN HYDROXYLAPATITE CALCIUM IRON BORON COPPER

MANGANESE BREAKING STRENGTH/

3/9,K/7 (Item 7 from file: 5)  
DIALOG(R) File 5:Biosis Previews(R)  
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0003895941 BIOSIS NO.: 198375079884  
**PURIFICATION AND PROPERTIES OF HYALURONIDASE EC-3.2.1.35 FROM HUMAN LIVER DIFFERENCES FROM AND SIMILARITIES TO THE TESTICULAR ENZYME**  
AUTHOR: GOLD E W (Reprint)  
AUTHOR ADDRESS: RESEARCH LAB, OHIO STATE UNIV COLL OF MED, COLUMBUS, OHIO 43210, USA\*\*USA  
JOURNAL: Biochemical Journal 205 (1): p69-74 1982  
ISSN: 0264-6021  
DOCUMENT TYPE: Article  
RECORD TYPE: Abstract  
LANGUAGE: ENGLISH

**ABSTRACT:** Human liver hyaluronidase was purified to homogeneity by (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> fractionation, chromatography on hydroxyapatite and DEAE-cellulose, and preparative disc polyacrylamide-gel electrophoresis. The enzyme had a pH optimum of 3.8-4.0, a MW (determined by gel filtration) of 76,000, and a Km of 0.05 mg/ml for purified human umbilical-cord hyaluronic acid. It generally resembled hyaluronidases studied in other tissues which are believed to be lysosomal, but shared a number of characteristics with a partially purified bovine testicular hyaluronidase. Neither enzyme exhibited inhibition by high concentrations of substrate, but both were competitively inhibited by dermatan sulfate and keratin sulfate. Both enzymes exhibited increased activity in the presence of albumin, probably owing to an increased susceptibility of substrate to enzyme action. The liver enzyme was inhibited by NaCl, but the testicular enzyme exhibited an increase in activity in the presence of the salt which was similar to the effect observed with albumin. The different response toward Cl<sup>-</sup> ion appeared to be the most significant difference between the 2 enzymes.

REGISTRY NUMBERS: 37326-33-3: EC-3.2.1.35; 16887-00-6: CHLORIDE ION  
DESCRIPTORS: BOVINE CHLORIDE ION

DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Enzymology-- Biochemistry and Molecular Biophysics

BIOSYSTEMATIC NAMES: Bovidae--Artiodactyla, Mammalia, Vertebrata, Chordata, Animalia; Hominidae--Primates, Mammalia, Vertebrata, Chordata, Animalia

COMMON TAXONOMIC TERMS: Artiodactyls; Nonhuman Vertebrates; Nonhuman Mammals; Animals; Chordates; Humans; Mammals; Primates; Vertebrates

CHEMICALS & BIOCHEMICALS: EC-3.2.1.35; CHLORIDE ION

CONCEPT CODES:

02508 Cytology - Human

10010 Comparative biochemistry

10064 Biochemistry studies - Proteins, peptides and amino acids

10069 Biochemistry studies - Minerals

10504 Biophysics - Methods and techniques

10802 Enzymes - General and comparative studies: coenzymes

10806 Enzymes - Chemical and physical

12100 Movement

14004 Digestive system - Physiology and biochemistry  
16504 Reproductive system - Physiology and biochemistry  
25502 Development and Embryology - General and descriptive

BIOSYSTEMATIC CODES:

85715 Bovidae  
86215 Hominidae

ABSTRACT: Human liver hyaluronidase was purified to homogeneity by (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> fractionation, chromatography on hydroxyapatite and DEAE-cellulose, and preparative disc polyacrylamide-gel electrophoresis. The enzyme had a pH optimum...

...inhibition by high concentrations of substrate, but both were competitively inhibited by dermatan sulfate and keratin sulfate. Both enzymes exhibited increased activity in the presence of albumin, probably owing to an...

3/9,K/8 (Item 1 from file: 24)  
DIALOG(R) File 24:CSA Life Sciences Abstracts  
(c) 2006 CSA. All rts. reserv.

0001974108 IP ACCESSION NO: 4515338  
Identification of tuftelin- and amelogenin-interacting proteins using the yeast two-hybrid system

Paine, CT; Paine, ML; Snead, ML  
University of Southern California, School of Dentistry, Center for Craniofacial Molecular Biology, 2250 Alcazar Street, Los Angeles, California 90033, USA

Connective Tissue Research, v 38, n 1-4, p 257-267, 1998  
PUBLICATION DATE: 1998

CONFERENCE:

6. Int. Symp. on the Composition, Properties and Fundamental Structure of Tooth Enamel, Lake Arrowhead, CA (USA), 11-15 May 1997

DOCUMENT TYPE: Journal Article; Conference

RECORD TYPE: Abstract

LANGUAGE: English

SUMMARY LANGUAGE: English

ISSN: 0300-8207

FILE SEGMENT: Calcium & Calcified Tissue Abstracts

ABSTRACT:

Biomineralization of enamel is a complex process that involves the eventual replacement of an extracellular protein matrix by hydroxyapatite crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin and ameloblastin. Assembly of the enamel extracellular matrix from these component proteins is believed to be critical in producing a matrix competent to undergo mineral replacement. Enamel formation is a complex process and additional proteins are likely to have a role in the assembly of the extracellular matrix. In order to identify additional proteins involved in the assembly process, the yeast two-hybrid system developed by Fields and Song (1989) has been implemented. This system allows for the identification of unknown proteins

that interact with proteins of interest. Typically a known protein is used as "bait" to screen a cDNA expression library of interest. In our studies, tuftelin or amelogenin have been used to screen a mouse tooth library produced from one day old pups. A library screening of six million clones with amelogenin as bait resulted in eleven positive clones all of which show high homology to the human leukocyte antigen-B (HLA-B) associated transcript (BAT) family of genes. A library screening of one million clones using tuftelin as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either keratin K5 or keratin K6, four are constitutively expressed and the remaining seven are novel. Further characterization of the proteins shown to interact with amelogenin or tuftelin may shed additional light on this complex process of enamel matrix assembly.

DESCRIPTORS: Teeth; Dental enamel; Keratin ; Mineralization; Extracellular matrix; Crystallization; Hydroxyapatite ; tuftelin; amelogenin; yeast two-hybrid system  
SUBJ CATG: 20079, Teeth

ABSTRACT:

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DESCRIPTORS: Teeth; Dental enamel; Keratin ; Mineralization; Extracellular matrix; Crystallization; Hydroxyapatite ; tuftelin; amelogenin; yeast two-hybrid system

3/9,K/9 (Item 1 from file: 34)

DIALOG(R)File 34:SciSearch(R) Cited Ref Sci  
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15540807 Genuine Article#: 081YL Number of References: 33

Title: Structure of white rhinoceros (*Ceratotherium simum*) horn investigated by X-ray histology with implications computed tomography and for growth and external form

Author(s): Hieronymus TL (REPRINT) ; Witmer LM; Ridgely RC

Corporate Source: Ohio Univ,Dept Sci Biol,Irvine Hall/Athens//OH/45701 (REPRINT); Ohio Univ,Dept Sci Biol,Athens//OH/45701; Ohio Univ,Coll Osteopath Med, Dept Biomed Sci,Athens//OH/45701(Th108702@ohiou.edu)

Journal: JOURNAL OF MORPHOLOGY, 2006, V267, N10 (OCT), P1172-1176

ISSN: 0362-2525 Publication date: 20061000

Publisher: WILEY-LISS, DIV JOHN WILEY & SONS INC, 111 RIVER ST, HOBOKEN, NJ 07030 USA

Language: English Document Type: ARTICLE

Geographic Location: USA

Journal Subject Category: ANATOMY & MORPHOLOGY

Abstract: The nasal and frontal horns of two individuals of *Ceratotherium simum* were examined by x-ray computed tomography (CT scanning), gross observation of sectioned horn, and light microscopy of histological

sections of the horn tissue. CT scans of both sets of horns reveal a periodic banding pattern that is evident upon gross observation of sections as darker bands of tissue. The overlap of these bands in both histological and CT slices suggests the presence of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely hydroxyapatite or octocalcium phosphate). The distribution of these two components in the horns is hypothesized to contribute to the differential wear patterns that produce the characteristic sweeping conical shape of rhinoceros horn from what otherwise (in the absence of wear and UV exposure) would be cylindrical blocks of constantly growing cornified papillary epidermis. Although extant rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of keratin-and-bone horns such as those of bovid artiodactyls, the tissue structures that make up the horn are strikingly convergent with other examples of papillary cornified epidermis found in horses, artiodactyls, cetaceans, and birds.

Descriptors--Author Keywords: anatomy ; histology ; tomography ; Ceratotherium ; rhinoceros ; integument ; keratin ; horn

Identifiers--KeyWord Plus(R): MECHANICAL-PROPERTIES; FRACTURE-TOUGHNESS; FEATHER KERATIN ; HOOF KERATIN ; DESIGN; PHASE; BILL; BONE

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...Abstract: of both a photoabsorbent component (melanin) and a radiodense

component (calcium phosphate salts, most likely hydroxyapatite or octocalcium phosphate). The distribution of these two components in the horns is hypothesized to...

...rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of keratin -and-bone horns such as those of bovid artiodactyls, the tissue structures that make up...

...Identifiers--MECHANICAL-PROPERTIES; FRACTURE-TOUGHNESS; FEATHER KERATIN ; HOOF KERATIN ; DESIGN; PHASE; BILL; BONE

3/9,K/10 (Item 2 from file: 34)  
DIALOG(R) File 34:SciSearch(R) Cited Ref Sci  
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14609630 Genuine Article#: 988SL Number of References: 62  
**Title: Applications of X-ray powder diffraction in materials chemistry**  
Author(s): Skakle J (REPRINT)  
Corporate Source: Univ Aberdeen,Dept Chem,Meston Walk/Aberdeen AB24  
3UE//Scotland/ (REPRINT); Univ Aberdeen,Dept Chem,Aberdeen AB24  
3UE//Scotland/(j.skakle@abdn.ac.uk)  
Journal: CHEMICAL RECORD, 2005, V5, N5, P252-262  
ISSN: 1527-8999 Publication date: 20050000  
Publisher: JOHN WILEY & SONS INC, 111 RIVER ST, HOBOKEN, NJ 07030 USA  
Language: English Document Type: ARTICLE  
Geographic Location: Scotland  
Journal Subject Category: CHEMISTRY, MULTIDISCIPLINARY  
Abstract: X-ray powder diffraction is a standard technique in materials chemistry, yet it is often still used in the laboratory as a "one-hit" technique, e.g. for fingerprinting and following the progress of reactions. It is important, however, that the wealth of information available from powder data is not overlooked. While it is only possible here to scratch the surface of possibilities, a range of examples from our research is used to emphasize some of the more accessible techniques and to highlight successes as well as potential problems. The first example is the study of solid solution formation in the oxide systems Ba<sub>3-3x</sub>La<sub>2x</sub>V<sub>208</sub> and Sr<sub>4-x</sub>BaxMn<sub>3010</sub> and in the silicate-hydroxyapatite bioceramic, Ca-10(PO<sub>4</sub>)<sub>(6-x)</sub>(SiO<sub>4</sub>)<sub>(x)</sub>(OH)<sub>(2-x)</sub>. Database mining is also explored, using three phases within the pseudobinary phase diagram Li<sub>3</sub>SbO<sub>4</sub>-CuO as examples. All three phases presented different challenges: the structure of Li<sub>3</sub>SbO<sub>4</sub> had been previously reported in higher symmetry than was actually the case, Li<sub>3</sub>Cu<sub>2</sub>SbO<sub>6</sub> was found to be isostructural with Li<sub>2</sub>TiO<sub>3</sub> but the cation ordering had to be rationalized, and Li<sub>3</sub>CuSbO<sub>5</sub> was believed to be triclinic, presenting challenges in indexing the powder pattern. Quantitative phase analysis is briefly discussed, with the emphasis both on success (determination of amorphous phase content in a novel cadmium arsenate phase) and on possible failure (compositional analysis in bone mineral); the reasons for the problems in the latter are also explored. Finally, the use of an area detector system has been shown to be of value in the study of orientational effects (or lack of them) in non- and partially-ordered biomaterials, including p-HEMA, annulus fibrosis of lumbar discs, and keratin in the horn of cow's hooves. (c) 2005 The Japan Chemical Journal Forum and Wiley Periodicals, Inc.

Descriptors--Author Keywords: x-ray diffraction ; solid state structures ; analytical methods

Identifiers--KeyWord Plus(R): CRYSTAL-STRUCTURE DETERMINATION; GENETIC

ALGORITHM; DAIRY-CATTLE; RATIO; HYDROXYAPATITE; PREDICTION; LAMENESS;  
COLLAGEN; LI3SBO4; PHASES

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...Abstract: solution formation in the oxide systems Ba<sub>3-3x</sub>La<sub>2x</sub>V<sub>2</sub>O<sub>8</sub> and Sr<sub>4-x</sub>Ba<sub>x</sub>Mn<sub>3</sub>O<sub>10</sub> and in the silicate- **hydroxyapatite** bioceramic, Ca-10(PO<sub>4</sub>)<sub>(6-x)</sub>(SiO<sub>4</sub>)<sub>(x)</sub>(OH)<sub>(2-x)</sub>. Database mining is also...

...in non- and partially-ordered biomaterials, including p-HEMA, annulus fibrosis of lumbar discs, and keratin in the horn of cow's hooves.

(c) 2005 The Japan Chemical Journal Forum and...

...Identifiers--CRYSTAL-STRUCTURE DETERMINATION; GENETIC ALGORITHM; DAIRY-CATTLE; RATIO; **HYDROXYAPATITE**; PREDICTION; LAMENESS; COLLAGEN; Li<sub>3</sub>SB<sub>4</sub>; PHASES

? t s3/9,k/11-21

3/9,K/11 (Item 3 from file: 34)

DIALOG(R) File 34:SciSearch(R) Cited Ref Sci

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13190177 . Genuine Article#: 855ZW Number of References: 17

Title: Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation

Author(s): Tachibana A; Kaneko S; Tanabe T; Yamauchi K (REPRINT)

Corporate Source: Osaka City Univ,Grad Sch Engn, Dept Appl & Bioappl Chem, Sumiyoshi Ku,Sugimoto 3-3-138/Osaka 5588585//Japan/ (REPRINT); Osaka City Univ,Grad Sch Engn, Dept Appl & Bioappl Chem, Sumiyoshi Ku,Osaka 5588585//Japan/(tatibana@biao.eng.osaka-cu.ac.jp; Yamauchi@biao.eng.osaka-cu.ac.jp)

Journal: BIOMATERIALS, 2005, V26, N3 (JAN), P297-302

ISSN: 0142-9612 Publication date: 20050100

Publisher: ELSEVIER SCI LTD, THE BOULEVARD, LANGFORD LANE, KIDLINGTON, OXFORD OX5 1GB, OXON, ENGLAND

Language: English Document Type: ARTICLE

Geographic Location: Japan

Journal Subject Category: ENGINEERING, BIOMEDICAL; MATERIALS SCIENCE, BIOMATERIALS

Abstract: Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium phosphate hybrid biomaterials were described. Firstly, the CaP-precipitated sponges were obtained by only the immersion of the carboxyl-sponges, chemically introduced with high amount of carboxyl groups on the sponges, in calcium and phosphate ions containing buffers such as PBS(+) for only 1-3 days. Neither sponge, introduced with amino or amido groups or non-treated, gave significant calcium phosphate precipitation. The carboxyl-sponges were mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the **hydroxyapatite** particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped **hydroxyapatite** particles might interact with keratin protein of the

sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast differentiation marker, suggested that both hybrid sponges, CaP-precipitated and trapped sponges, alter the differentiation pattern of preosteoblasts, MC3T3-E1. (C) 2004 Elsevier Ltd. All rights reserved.

Descriptors--Author Keywords: wool keratin sponge ; hydroxyapatite hybrid ; osteoblast scaffolds ; osteoblast differentiation

Identifiers--KeyWord Plus(R): BONE-LIKE APATITE; BIOMATERIAL; LYSOZYME; COLLAGEN; PROTEIN; GROWTH; ASSAY; CELLS

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YAMAUCHI K, 1997, V9, P117, COLLOID SURFACE B

Title: Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation

Abstract: Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium...

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3/9,K/12 (Item 4 from file: 34)

DIALOG(R) File 34:SciSearch(R) Cited Ref Sci  
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13112099. Genuine Article#: 849ZP Number of References: 28

Title: Characterization of Prismalin-14, a novel matrix protein from the prismatic layer of the Japanese pearl oyster (*Pinctada fucata*)

Author(s): Suzuki M; Murayama E; Inoue H; Ozaki N; Tohse H; Kogure T; Nagasawa H (REPRINT)

Corporate Source: Univ Tokyo,Dept Appl Biol Chem, Grad Sch Agr & Life Sci, Bunkyo Ku, 1-1-1 Yayoi/Tokyo 1138657//Japan/ (REPRINT); Univ Tokyo,Dept Appl Biol Chem, Grad Sch Agr & Life Sci, Bunkyo Ku,Tokyo 1138657//Japan/; Japan Sci & Technol Agcy,CREST,Saitama 3320012//Japan/

; Univ Tokyo,Dept Earth & Planetary Sci, Grad Sch Sci, Bunkyo Ku,Tokyo  
1130033//Japan/(anagahi@mail.ecc.u-tokyo.ac.jp)

Journal: BIOCHEMICAL JOURNAL, 2004, V382, 1 (AUG 15), P205-213

ISSN: 0264-6021 Publication date: 20040815

Publisher: PORTLAND PRESS, 59 PORTLAND PLACE, LONDON W1N 3AJ, ENGLAND

Language: English Document Type: ARTICLE

Geographic Location: Japan

Journal Subject Category: BIOCHEMISTRY & MOLECULAR BIOLOGY

Abstract: The mollusc shell is a hard tissue consisting of calcium carbonate and organic matrices. The organic matrices are believed to play important roles in shell formation. In the present study, we extracted and purified a novel matrix protein, named Prismalin-14, from the acid-insoluble fraction of the prismatic layer of the shell of the Japanese pearl oyster (*Pinctada fucata*), and determined its whole amino acid sequence by a combination of amino acid sequence analysis and MS analysis of the intact protein and its enzymic digests. Prismalin-14 consisted of 105 amino acid residues, including PIYR repeats, a Gly/Tyr-rich region and N- and C-terminal Asp-rich regions. Prismalin-14 showed inhibitory activity on calcium carbonate precipitation and calcium-binding activity in vitro. The scanning electron microscopy images revealed that Prismalin-14 affected the crystallization of calcium carbonate in vitro. A cDNA encoding Prismalin-14 was cloned and its expression was analysed. The amino acid sequence deduced from the nucleotide sequence of Prismalin-14 cDNA was identical with that determined by peptide sequencing. Northern-blot analysis showed that a Prismalin-14 mRNA was expressed only at the mantle edge. In situ hybridization demonstrated that a Prismalin-14 mRNA was expressed strongly in the inner side of the outer fold of the mantle. These results suggest that Prismalin-14 is a framework protein that plays an important role in the regulation of calcification of the prismatic layer of the shell.

Descriptors--Author Keywords: biomineralization ; calcification ; matrix protein ; mollusc shell ; pearl oyster ; prismatic layer

Identifiers--KeyWord Plus(R): ANTI-CALCIFICATION; MOLECULAR-CLONING; NACREOUS LAYER; BINDING; SHELL; EXPRESSION; KERATIN; CDNA; HYDROXYAPATITE; CRAYFISH

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...Identifiers--ANTI-CALCIFICATION; MOLECULAR-CLONING; NACREOUS LAYER;  
BINDING; SHELL; EXPRESSION; KERATIN; CDNA; HYDROXYAPATITE; CRAYFISH

3/9,K/13 (Item 5 from file: 34)

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12896914 Genuine Article#: 829WK Number of References: 95

Title: The mechanical efficiency of natural materials

Author(s): Wegst UGK; Ashby MF (REPRINT)

Corporate Source: Univ Cambridge,Dept Engn, Engn Design Ctr,Trumpington St/Cambridge CB2 1PZ//England/ (REPRINT); Univ Cambridge,Dept Engn, Engn Design Ctr,Cambridge CB2 1PZ//England/; Max Planck Inst Met Res,D-70569 Stuttgart//Germany/ (mfaz@eng.cam.ac.uk)

Journal: PHILOSOPHICAL MAGAZINE, 2004, V84, N21 (JUL 21), P2167-2181

ISSN: 1478-6443 Publication date: 20040721

Publisher: TAYLOR & FRANCIS LTD, 4 PARK SQUARE, MILTON PARK, ABINGDON OX14 4RN, OXON, ENGLAND

Language: English Document Type: REVIEW

Geographic Location: England; Germany

Journal Subject Category: MATERIALS SCIENCE, MULTIDISCIPLINARY; MECHANICS; METALLURGY & METALLURGICAL ENGINEERING; PHYSICS, APPLIED; PHYSICS, CONDENSED MATTER

Abstract: The materials of nature, for example cellulose, lignin, keratin, chitin, collagen and hydroxyapatite, and the structures made from them, for example bamboo, wood, antler and bone, have a remarkable range of mechanical properties. These can be compared by presenting them as material property charts, well known for the materials of engineering. Material indices (significant combinations of properties) can be plotted on to the charts, identifying materials with extreme values of an index, suggesting that they have evolved to carry particular modes of loading, or to sustain large tensile or flexural deformations, without failure. This paper describes a major revision and update of a set of property charts for natural material published some 8 years ago by Ashby et al. with examples of their use to study mechanical efficiency in nature.

Identifiers--KeyWord Plus(R): WALLABY TAIL TENDONS; TRABECULAR BONE; CONSTITUTIVE-EQUATIONS; FRACTURE PROPERTIES; MAMMALIAN TENDONS;

PATELLAR TENDON; CANCELLOUS BONE; MACADAMIA NUTS; CORAL SKELETON; BEHAVIOR

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**Abstract:** The materials of nature, for example cellulose, lignin, keratin , chitin, collagen and hydroxyapatite , and the structures made from them, for example bamboo, wood, antler and bone, have a...

3/9,K/14 (Item 6 from file: 34)  
DIALOG(R) File 34:SciSearch(R) Cited Ref Sci  
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10435091 Genuine Article#: 528XF Number of References: 41  
**Title: Bone tissue fluorescence reduction for visible laser Raman spectroscopy**  
Author(s): Shea DA; Morris MD (REPRINT)  
Corporate Source: Univ Michigan,Dept Chem,Ann Arbor//MI/48109 (REPRINT);  
Univ Michigan,Dept Chem,Ann Arbor//MI/48109  
Journal: APPLIED SPECTROSCOPY, 2002, V56, N2 (FEB), P182-186  
ISSN: 0003-7028 Publication date: 20020200  
Publisher: SOC APPLIED SPECTROSCOPY, 201B BROADWAY ST, FREDERICK, MD 21701 USA

Language: English Document Type: ARTICLE  
Geographic Location: USA  
Journal Subject Category: INSTRUMENTS & INSTRUMENTATION; SPECTROSCOPY  
**Abstract:** We report the successful reduction of background fluorescence in bone tissue by photo-irradiation with green laser light. Irradiation of

bone tissue with intense green light has been shown to be non-destructive and to reduce permanently over 70% of the fluorescence background. The laser power dependence of the fluorescence reduction was found to be nonlinear.

Descriptors--Author Keywords: Raman spectroscopy ; bone ; photobleaching ; principal components analysis ; factor analysis

Identifiers--KeyWord Plus(R): UV-RADIATION; COLLAGEN; HYDROXYAPATITE; PHOTODEGRADATION; SPECTRA; KERATIN; CELLS

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...Identifiers--UV-RADIATION; COLLAGEN; HYDROXYAPATITE; PHOTODEGRADATION; SPECTRA; KERATIN; CELLS

3/9,K/15 (Item 7 from file: 34)

DIALOG(R) File 34:SciSearch(R) Cited Ref Sci

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07282281 Genuine Article#: 145WF Number of References: 56

**Title:** Identification of tuftelin- and amelogenin-interacting proteins using the yeast two-hybrid system

**Author(s):** Paine CT (REPRINT) ; Paine ML; Snead ML

**Corporate Source:** UNIV SO CALIF,CTR CRANIOFACIAL MOL BIOL, SCH DENT, 2250 ALCAZAR ST/LOS ANGELES//CA/90033 (REPRINT)

**Journal:** CONNECTIVE TISSUE RESEARCH, 1998, V39, N1-3, P257-267

**ISSN:** 0300-8207 **Publication date:** 19980000

**Publisher:** GORDON BREACH SCI PUBL LTD, C/O STBS LTD, PO BOX 90, READING RG1 8JL, BERKS, ENGLAND

**Language:** English **Document Type:** ARTICLE

**Geographic Location:** USA

**Subfile:** CC LIFE--Current Contents, Life Sciences;

**Journal Subject Category:** ORTHOPEDICS; CELL BIOLOGY

**Abstract:** Biominerization of enamel is a complex process that involves the eventual replacement of an extracellular protein matrix by hydroxyapatite crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin and ameloblastin. Assembly of the enamel extracellular matrix from these component proteins is believed to be critical in producing a matrix competent to undergo mineral replacement. Enamel formation is a complex process and additional proteins are likely to have a role in the assembly of the extracellular matrix. In order to identify additional proteins involved in the assembly process, the yeast two-hybrid system developed by Fields and Song (1989) has been implemented. This system allows for the identification of unknown proteins that interact with proteins of interest. Typically a known protein is used as 'bait' to screen a cDNA expression library of interest. In our studies, tuftelin or amelogenin have been used to screen a mouse teeth library produced from one day old pups. A library screening of six million clones with amelogenin as bait resulted in eleven positive clones all of which show high homology to the human leukocyte antigen-B (HLA-B) associated transcript (BAT) family of genes. A library screening of one million clones using tuftelin as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either keratin K5 or keratin K6, four are constitutively expressed and the remaining seven are novel. Further characterization of the proteins shown to interact with amelogenin or tuftelin may shed additional light on this complex process of enamel matrix assembly.

**Descriptors--Author Keywords:** biominerization ; enamel cDNA expression library ; self-assembly and odontogenesis

**Identifiers--KeyWord Plus(R):** ENAMELIN TUFTELIN; TOOTH DEVELOPMENT; GENE-EXPRESSION; DNA-POLYMERASE; MOUSE; SEQUENCE; MATRIX; DIFFERENTIATION; ORGANOGENESIS; LOCALIZATION

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...Abstract: is a complex process that involves the eventual replacement of an extracellular protein matrix by hydroxyapatite crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin...

...as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either keratin K5 or keratin K6, four are constitutively expressed and the remaining seven are novel. Further characterization of the...

3/9,K/16 (Item 8 from file: 34)  
DIALOG(R) File 34:SciSearch(R) Cited Ref Sci  
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05198600 Genuine Article#: VG397 Number of References: 47  
**Title: THE LOCALIZATION OF EPITHELIAL ROOT SHEATH-CELLS DURING CEMENTUM FORMATION IN RAT MOLARS**

Author(s): ALATLI I; LUNDMARK C; HAMMARSTROM L

Corporate Source: KAROLINSKA INST,CTR ORAL BIOL,NOVUM,BOX 4064/S-14104  
HUDDINGE//SWEDEN/

Journal: JOURNAL OF PERIODONTAL RESEARCH, 1996, V31, N6 (AUG), P433-440  
ISSN: 0022-3484

Language: ENGLISH Document Type: ARTICLE

Geographic Location: SWEDEN

Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences

Journal Subject Category: DENTISTRY/ORAL SURGERY & MEDICINE

**Abstract:** The purpose of this study was to investigate the distribution of epithelial cells and the fate of the basement membrane along the root surface of rat molars during cementogenesis, and to test the hypothesis that the Hertwig's epithelial root sheath (HERS) cells remain on the root surface if mineralization is inhibited. To demonstrate the HERS cells and basement membrane, immunohistochemistry with antibodies against keratin and laminin were used. The dentin matrix mineralization was inhibited by a single injection of 1-hydroxyethylidene-1,1-bisphosphonate (HEBP). A modified Gomori staining method was used to monitor the inhibition of mineral formation in dentin and cementum. Paraffin sections were stained with haematoxylin-eosin, and freeze-dried sections were used for Gomori and immunohistochemical stainings. We found that the formation of acellular cementum was suppressed above the dentin with inhibited mineralization. Instead, a hyperplastic matrix, different from acellular cementum, covered the dentin. This hyperplastic cementum had keratin - and laminin-positive cells incorporated; such cells were never incorporated in normal acellular cementum. The later formation of cellular cementum correlated, in controls, with the disappearance of HERS cells from the root surface. Treatment with HEBP resulted in a persistent presence of epithelial cells, interpreted as an inhibition of their disappearance. In conclusion, there is evidence that the cells of HERS are involved in the development of both acellular and cellular cementum. The developmental processes of these tissues appear in some way to be influenced by or associated with the initial mineralization of the dentin.

**Descriptors--Author Keywords:** CEMENTUM ; HERS ; KERATIN ; LAMININ ; HEBP  
**Identifiers--KeyWords Plus:** INTERMEDIATE CEMENTUM; BASEMENT-MEMBRANE;

DENTIN; MOUSE; HYDROXYAPATITE; PERIODONTIUM; LAMININ; HERTWIG;  
ENAMEL; MONKEY

**Research Fronts:** 94-4329 001 (K12 KERATIN IN RABBIT CORNEAL LIMBAL EPITHELIAL-CELLS; CYTOKERATIN EXPRESSION; INTERMEDIATE FILAMENT PROTEIN; DIAGNOSIS OF EPIDERMOLYSIS-BULLOSA SIMPLEX)

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...Abstract: mineralization is inhibited. To demonstrate the HERs cells and basement membrane, immunohistochemistry with antibodies against keratin and laminin were used. The dentin matrix mineralization was inhibited by a single injection of...

...Instead, a hyperplastic matrix, different from acellular cementum, covered the dentin. This hyperplastic cementum had keratin - and laminin-positive cells incorporated; such cells were never incorporated in normal acellular cementum. The...

...Identifiers--INTERMEDIATE CEMENTUM; BASEMENT-MEMBRANE; DENTIN; MOUSE; HYDROXYAPATITE; PERIODONTIUM; LAMININ; HERTWIG; ENAMEL; MONKEY

Research Fronts: 94-4329 001 (K12 KERATIN IN RABBIT CORNEAL LIMBAL EPITHELIAL-CELLS; CYTOKERATIN EXPRESSION; INTERMEDIATE FILAMENT PROTEIN; DIAGNOSIS OF EPIDERMOLYSIS-BULLOSA...)

3/9,K/17 (Item 9 from file: 34)  
DIALOG(R) File 34:SciSearch(R) Cited Ref Sci  
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03775581 Genuine Article#: QE293 Number of References: 50  
**Title: ATOMIC-FORCE MICROSCOPY OF THE MORPHOLOGY OF THE MATRIX AND MINERAL COMPONENTS OF THE OTOLITH OF HYPEROGLYPHE ANTARCTICA**

**Author(s): GAULDIE RW; XHIE J**  
**Corporate Source: UNIV HAWAII,SCH OCEAN & EARTH SCI & TECHNOL,HAWAII INST GEOPHYS & PLANETOL/HONOLULU//HI/96822**

**Journal: JOURNAL OF MORPHOLOGY, 1995, V223, N2 (FEB), P203-214**

**ISSN: 0362-2525**

**Language: ENGLISH Document Type: ARTICLE**

**Geographic Location: USA**

**Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences**

**Journal Subject Category: ANATOMY & MORPHOLOGY**

**Abstract:** The sagittal otolith of Hyperoglyphe antarctica (Centrolophidae: Teleostei) has a prismatic structure in which the anti-sulcal growth axes of each prism consist of a series of nested cones each composed of a mineral layer followed by an organic matrix layer. Broken sections show the mineral layers to be composed of stacks of crystals. Otolith matrix that has been decalcified and air-dried, or critical-point-dried, retains a periodic structure of repeating high and low matrix density. At high magnifications, both broken whole crystal surfaces and decalcified matrix surfaces have a granular structure. Chlorox-bleached whole otoliths also show a granular crystalline structure. At higher magnifications, the air-dried matrix showed a parallel fiber structure with similar dimensions to keratin fibers. (C) 1995 Wiley-Liss, Inc..

**Identifiers--KeyWords Plus: FISH OTOLITHS; FUNDULUS-HETEROCRITUS; ELECTRON-MICROSCOPE; CALCITE; GROWTH; ULTRASTRUCTURE; TEMPERATURE; ARAGONITE; OTOCONIA; RINGS**

**Research Fronts: 93-3649 002 (OTOLITH MICROSTRUCTURE; DAILY GROWTH INCREMENTS; FISH AGE VALIDATION; EARLY-LIFE HISTORY EVENTS)**

93-0242 001 (CHIRAL STATIONARY PHASES; OPTICAL RESOLUTION; HIGH-PERFORMANCE LIQUID-CHROMATOGRAPHY)

93-0486 001 (ATOMIC FORCE MICROSCOPY; IMAGING SURFACES; RECONSTITUTED BIOLOGICAL CHANNELS AT MOLECULAR RESOLUTION)

93-5794 001 (MIXED MONOLAYERS; ICE NUCLEATION; BIOMIMETIC MATERIALS CHEMISTRY; GELATINOUS MEMBRANE; ORGANIC TEMPLATE)

93-6422 001 (HYDROXYAPATITE INDUCTION; INVITRO BEHAVIOR; MACROPOROUS CALCIUM-PHOSPHATE CERAMICS)

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...Abstract: higher magnifications, the air-dried matrix showed a parallel fiber structure with similar dimensions to keratin fibers. (C) 1995 Wiley-Liss, Inc.

...Research Fronts: 001 (MIXED MONOLAYERS; ICE NUCLEATION; BIOMIMETIC MATERIALS CHEMISTRY; GELATINOUS MEMBRANE; ORGANIC TEMPLATE)  
93-6422 001 (HYDROXYAPATITE INDUCTION; INVITRO BEHAVIOR; MACROPOROUS CALCIUM-PHOSPHATE CERAMICS)

3/9,K/18 (Item 10 from file: 34)  
DIALOG(R) File 34:SciSearch(R) Cited Ref Sci  
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00826041 Genuine Article#: FA140 Number of References: 32  
Title: CONSTRUCTION OF A UNIFORM-ABUNDANCE (NORMALIZED) CDNA LIBRARY  
Author(s): PATANJALI SR; PARIMOO S; WEISSMAN SM  
Corporate Source: YALE UNIV,SCH MED,DEPT HUMAN GENET,333 CEDAR ST/NEW HAVEN//CT/06510  
Journal: PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED

STATES OF AMERICA, 1991, V88, N5, P1943-1947  
Language: ENGLISH Document Type: ARTICLE  
Geographic Location: USA  
Subfile: SciSearch; CC LIFE--Current Contents, Life Sciences  
Journal Subject Category: MULTIDISCIPLINARY SCIENCES  
Abstract: We have used a kinetic approach to construct cDNA libraries containing approximately equal representations of all sequences in a preparation of poly(A)+ RNA. Randomly primed cDNA fragments of a selected size range were cloned in lambda-phage vector. Inserts were amplified by the polymerase chain reaction (PCR), denatured, and self-annealed under optimized conditions. After extensive but incomplete reannealing, the single-stranded fraction was relatively depleted of more abundant species of cDNA. Libraries of these fragments are suitable for cDNA subtraction, screening, or selection by hybridization and make it possible to detect and analyze cDNA corresponding to species of mRNA present at a low level in a small fraction of the cells in a complex tissue.

Descriptors--Author Keywords: REASSOCIATION; HYBRIDIZATION; HYDROXYAPATITE

Identifiers--KeyWords Plus: RIBOSOMAL DNA SEGMENTS; MOLECULAR ANALYSIS; MESSENGER-RNA; HUMAN GENOME; SEQUENCES; GENES; ACID; EXTRACTION; FAMILY Research Fronts: 89-1447 002 (DEVELOPMENTALLY REGULATED GENE; CAPPING PROTEIN; CDNA SEQUENCE; GENOME ORGANIZATION)

89-3817 002 (POLYMERASE CHAIN-REACTION; AMPLIFIED GENOMIC DNA; DIRECT SEQUENCING)

89-0025 001 (CHLOROPHYLL FLUORESCENCE; PHOTOSYSTEM-II IN PEA LEAVES; DNA-DNA HYBRIDIZATION; MOLECULAR CLOCKS; NON-PHOTOCHEMICAL QUENCHING; THERMAL TOLERANCE)

89-0580 001 (HUMAN IMMUNODEFICIENCY VIRUS; HIV INFECTION; RECOMBINANT SOLUBLE CD4 RECEPTOR; PROTEIN EXPRESSION VIA A CIS-ACTING SEQUENCE)

89-1700 001 (INSITU HYBRIDIZATION USING CHROMOSOME-SPECIFIC ALPHA-SATELLITE DNA; GENOMIC ORGANIZATION; AVIAN KERATIN GENES; HIGHLY REPETITIVE SEQUENCE)

89-1708 001 (T-CELL RECEPTOR DELTA-GENES IN HUMAN T-CELL LEUKEMIAS; ANALYSIS OF JUNCTIONAL DIVERSITY; IMPLICATIONS FOR THYMIC DIFFERENTIATION)

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WILSON GN, 1978, V75, P5367, P NATL ACAD SCI USA  
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...Research Fronts: SEQUENCE)  
89-1700 001 (INSITU HYBRIDIZATION USING CHROMOSOME-SPECIFIC  
ALPHA-SATELLITE DNA; GENOMIC ORGANIZATION; AVIAN KERATIN GENES;  
HIGHLY REPETITIVE SEQUENCE)  
89-1708 001 (T-CELL RECEPTOR DELTA-GENES IN HUMAN T...

3/9,K/19 (Item 1 from file: 45)  
DIALOG(R) File 45:EMCare  
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01500532 EMCare No: 38950713  
Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward  
osteoblast cultivation and differentiation  
Tachibana A.; Kaneko S.; Tanabe T.; Yamauchi K.  
K. Yamauchi, Dept. of Appl. and Bioapplied Chem., Graduate School of  
Engineering, Osaka City Univ., S., Osaka Japan  
AUTHOR EMAIL: Yamauchi@bioa.eng.osaka-cu.ac.jp  
Biomaterials ( BIOMATERIALS ) (United Kingdom) 2005, 26/3 (297-302)  
CODEN: BIMAD ISSN: 0142-9612  
PUBLISHER ITEM IDENTIFIER: S014296120400170X  
DOCUMENT TYPE: Journal ; Article  
LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH  
NUMBER OF REFERENCES: 17  
RECORD TYPE: Abstract

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium phosphate hybrid biomaterials were described. Firstly, the CaP-precipitated sponges were obtained by only the immersion of the carboxyl-sponges, chemically introduced with high amount of carboxyl groups on the sponges, in calcium and phosphate ions containing buffers such as PBS(+) for only 1-3 days. Neither sponge, introduced with amino or amido groups or non-treated, gave significant calcium phosphate precipitation. The carboxyl-sponges were mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast differentiation marker, suggested that both hybrid sponges, CaP-precipitated and trapped sponges, alter the differentiation pattern of preosteoblasts, MC3T3-E1. (c) 2004 Elsevier Ltd. All rights reserved.

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DESCRIPTORS:

\* keratin ; \* hydroxyapatite ; \*osteoblast; \*wool calcium phosphate; 4 carboxyglutamic acid; protein; alkaline phosphatase; marker; acid protein; biomaterial; carboxyl group; calcium; phosphate; precipitation; calcification; cell density; immersion

**Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation**

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium...

...mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast...

DESCRIPTORS:

\* keratin ; \* hydroxyapatite ; \*osteoblast; \*wool

3/9,K/20 (Item 1 from file: 65)

DIALOG(R) File 65:Inside Conferences

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05013684 INSIDE CONFERENCE ITEM ID: CN052238262

**13:20 II Y09 Fabrication of keratin - hydroxyapatite hybrid sponges**

Kaneko, S.; Tachibana, A.; Tanabe, T.; Yamauchi, K.

CONFERENCE: Macromolecules;; SPSJ-Symposium; 52nd

POLYMER PREPRINTS JAPAN -ENGLISH EDITION-, 2003; VOL 52; NO 2 P: E 1033

Society of Polymer Science, Japan,, 2003

LANGUAGE: English DOCUMENT TYPE: Conference Preprinted abstracts

CONFERENCE SPONSOR: Society of Polymer Science, Japan

CONFERENCE LOCATION: Yamaguchi, Japan 2003; Sep (200309)

BRITISH LIBRARY ITEM LOCATION: 6547.715300

DESCRIPTORS: Macromolecules; Polymer science; SPSJ

**13:20 II Y09 Fabrication of keratin - hydroxyapatite hybrid sponges**

3/9,K/21 (Item 1 from file: 71)

DIALOG(R) File 71:ELSEVIER BIOBASE

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03490768 2006275957

Structure of White Rhinoceros (*Ceratotherium simum*) horn investigated by X-ray computed tomography and histology with implications for growth and external form

Hieronymus T.L.; Witmer L.M.; Ridgely R.C.

ADDRESS: T.L. Hieronymus, Department of Biological Sciences, Irvine Hall, Ohio University, Athens, OH 45701, United States

EMAIL: Th108702@ohiou.edu  
Journal: Journal of Morphology, 267/10 (1172-1176), 2006, United States  
CODEN: JOMOA  
ISSN: 0362-2525 eISSN: 1097-4687  
DOCUMENT TYPE: Article  
LANGUAGES: English SUMMARY LANGUAGES: English  
NO. OF REFERENCES: 33

The nasal and frontal horns of two individuals of *Ceratotherium simum* were examined by x-ray computed tomography (CT scanning), gross observation of sectioned horn, and light microscopy of histological sections of the horn tissue. CT scans of both sets of horns reveal a periodic banding pattern that is evident upon gross observation of sections as darker bands of tissue. The overlap of these bands in both histological and CT slices suggests the presence of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely **hydroxyapatite** or **octocalcium phosphate**). The distribution of these two components in the horns is hypothesized to contribute to the differential wear patterns that produce the characteristic sweeping conical shape of rhinoceros horn from what otherwise (in the absence of wear and UV exposure) would be cylindrical blocks of constantly growing cornified papillary epidermis. Although extant rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of **keratin**-and-bone horns such as those of bovid artiodactyls, the tissue structures that make up the horn are strikingly convergent with other examples of papillary cornified epidermis found in horses, artiodactyls, cetaceans, and birds. (c) 2006 Wiley-Liss, Inc.

DESCRIPTORS:

Anatomy; Histology; Tomography; Ceratotherium; Rhinoceros; Integument; Keratin ; Horn

SPECIES DESCRIPTORS:

*Ceratotherium simum*; Rhinoceros; Ceratotherium; Aves; Bovidae; Artiodactyla ; Rhinocerotidae; Cetacea; Equidae; *Equus caballus*

CLASSIFICATION CODE AND DESCRIPTION:

99 - General

...of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely **hydroxyapatite** or **octocalcium phosphate**). The distribution of these two components in the horns is hypothesized to...

...rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of **keratin**-and-bone horns such as those of bovid artiodactyls, the tissue structures that make up...

DESCRIPTORS:

Anatomy; Histology; Tomography; Ceratotherium; Rhinoceros; Integument; Keratin ; Horn  
? t s3/9,k/22-32

3/9,K/22 (Item 2 from file: 71)  
DIALOG(R) File 71:ELSEVIER BIOBASE  
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01053759 1999030536 .

**Identification of tuftelin- and amelogenin-interacting proteins using the yeast two-hybrid system**

Paine C.T.; Paine M.L.; Snead M.L.

ADDRESS: C.T. Paine, University of Southern California, School of Dentistry, Ctr. for Craniofacial Molec. Biology, 2250 Alcazar Street, Los Angeles, CA 90033, United States

Journal: Connective Tissue Research, 38/1-4 (257-267), 1998, United Kingdom

CODEN: CVTRB

ISSN: 0300-8207

DOCUMENT TYPE: Conference Paper

LANGUAGES: English SUMMARY LANGUAGES: English

NO. OF REFERENCES: 56

Biomineralization of enamel is a complex process that involves the eventual replacement of an extracellular protein matrix by **hydroxyapatite** crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin and ameloblastin. Assembly of the **enamel** extracellular matrix from these component proteins is believed to be critical in producing a matrix competent to undergo mineral replacement. Enamel formation is a complex process and additional proteins are likely to have a role in the assembly of the extracellular matrix. In order to identify additional proteins involved in the assembly process, the yeast two-hybrid system developed by Fields and Song (1989) has been implemented. This system allows for the identification of unknown proteins that interact with proteins of interest. Typically a known protein is used as 'bait' to screen a cDNA expression library of interest. In our studies, tuftelin or amelogenin have been used to screen a mouse tooth library produced from one day old pups. A library screening of six million clones with amelogenin as bait resulted in eleven positive clones all of which show high homology to the human leukocyte antigen-B (HLA-B) associated transcript (BAT) family of genes. A library screening of one million clones using tuftelin as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either **keratin K5** or **keratin K6**, four are constitutively expressed and the remaining seven are novel. Further characterization of the proteins shown to interact with amelogenin or tuftelin may shed additional light on this complex process of enamel matrix assembly.

**DESCRIPTORS:**

Biomineralization; Enamel cDNA expression library; Self-assembly and odontogenesis

**CLASSIFICATION CODE AND DESCRIPTION:**

82.2.12.2 - PROTEIN BIOCHEMISTRY / STRUCTURAL STUDIES / Molecular Recognition / Protein-protein interaction

89.4.1.1 - CELL AND DEVELOPMENTAL BIOLOGY / EXTRACELLULAR MATRIX (STRUCTURE AND FUNCTION) / Extracellular Matrix / Structure and composition

...is a complex process that involves the eventual replacement of an extracellular protein matrix by **hydroxyapatite** crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin...

...as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either **keratin K5** or **keratin K6**, four are constitutively expressed and the remaining seven are novel. Further characterization of the...

3/9,K/23 (Item 1 from file: 73)

DIALOG(R) File 73:EMBASE

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12725738 EMBASE No: 2004309901

**Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation**

Tachibana A.; Kaneko S.; Tanabe T.; Yamauchi K.

K. Yamauchi, Dept. of Appl. and Bioapplied Chem., Graduate School of Engineering, Osaka City Univ., S., Osaka Japan

AUTHOR EMAIL: Yamauchi@bioa.eng.osaka-cu.ac.jp

Biomaterials ( BIOMATERIALS ) (United Kingdom) 2005, 26/3 (297-302)

CODEN: BIMAD ISSN: 0142-9612

PUBLISHER ITEM IDENTIFIER: S014296120400170X

DOCUMENT TYPE: Journal ; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 17

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium phosphate hybrid biomaterials were described. Firstly, the CaP-precipitated sponges were obtained by only the immersion of the carboxyl-sponges, chemically introduced with high amount of carboxyl groups on the sponges, in calcium and phosphate ions containing buffers such as PBS(+) for only 1-3 days. Neither sponge, introduced with amino or amido groups or non-treated, gave significant calcium phosphate precipitation. The carboxyl-sponges were mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast differentiation marker, suggested that both hybrid sponges, CaP-precipitated and trapped sponges, alter the differentiation pattern of preosteoblasts, MC3T3-E1. (c) 2004 Elsevier Ltd. All rights reserved.

DRUG DESCRIPTORS:

\* keratin ; \* hydroxyapatite

calcium phosphate; biomaterial; carboxyl group; calcium ion; phosphate; buffer; 4 carboxyglutamic acid; alkaline phosphatase

MEDICAL DESCRIPTORS:

\*osteoblast; \*cell differentiation

hybrid; precipitation; calcification; protein expression; cell line; nonhuman; mouse; controlled study; animal cell; article; priority journal

CAS REGISTRY NO.: 1306-06-5, 51198-94-8 ( hydroxyapatite ); 10103-46-5,

13767-12-9, 14358-97-5, 7758-87-4 (calcium phosphate); 14127-61-8 (

calcium ion); 14066-19-4, 14265-44-2 (phosphate); 53861-57-7 (4 carboxyglutamic acid); 9001-78-9 (alkaline phosphatase)

SECTION HEADINGS:

027 Biophysics, Bioengineering and Medical Instrumentation

029 Clinical and Experimental Biochemistry

033 Orthopedic Surgery

**Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward**

## **osteoblast cultivation and differentiation**

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium...

...mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast...

### **DRUG DESCRIPTORS:**

\* keratin ; \* hydroxyapatite

...CAS REGISTRY NO.: 51198-94-8 ( hydroxyapatite ); 10103-46-5...

3/9,K/24 (Item 2 from file: 73)

DIALOG(R) File 73:EMBASE

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12047050 EMBASE No: 2003158648

Implantable applications of chitin and chitosan

Khor E.; Lim L.Y.

E. Khor, Department of Chemistry, National University of Singapore, 3 Science Drive 3, Kent Ridge, Singapore 117543 Singapore

AUTHOR EMAIL: chmkhore@nus.edu.sg

Biomaterials ( BIOMATERIALS ) (United Kingdom) 2003, 24/13 (2339-2349)

CODEN: BIMAD ISSN: 0142-9612

DOCUMENT TYPE: Journal ; Article

LANGUAGE: ENGLISH SUMMARY LANGUAGE: ENGLISH

NUMBER OF REFERENCES: 91

Chitin, extracted primarily from shellfish sources, is a unique biopolymer based on the N-acetyl-glucosamine monomer. More than 40 years have lapsed since this biopolymer had aroused the interest of the scientific community around the world for its potential biomedical applications. Chitin, together with its variants, especially its deacetylated counterpart chitosan, has been shown to be useful as a wound dressing material, drug delivery vehicle and increasingly a candidate for tissue engineering. The promise for this biomaterial is vast and will continue to increase as the chemistry to extend its capabilities and new biomedical applications are investigated. It is interesting to note that a majority of this work has come from Asia. Japan has been the undisputed leader, but other Asian nations, namely Korea, Singapore, Taiwan and Thailand have also made notable contributions. More recently, China has joined the club to become an increasingly major research source for chitin and chitosan in Asia. This review surveys select works of key groups in Asia developing chitin and chitosan materials for implantable biomedical applications. (c) 2003 Elsevier Science Ltd. All rights reserved.

### **DRUG DESCRIPTORS:**

\*chitin--drug combination--cb; \*chitin--drug comparison--cm; \*chitin--pharmaceutics--pr; \*chitosan--pharmaceutics--pr

biomaterial--pharmaceutics--pr; hydroxyapatite ; calcium oxide; bone cement; calcium carbonate; gelatin; alginic acid--pharmaceutics--pr; n

acetylglucosamine; hyaluronic acid; keratin --pharmaceutics--pr; keratin --pharmacology--pd; macrogol--pharmaceutics--pr; sulfadiazine silver --pharmaceutics--pr; sulfadiazine silver--pharmacology--pd; chitin derivative; triamcinolone acetonide--drug therapy--dt; triamcinolone acetonide--pharmaceutics--pr; acrylic acid--pharmaceutics--pr; cyanocobalamin--pharmaceutics--pr; Salvia miltiorrhiza extract --pharmaceutics--pr; microsphere--pharmaceutics--pr; microsphere --intravenous drug administration--iv; polyglactin--drug combination--cb; polyglactin--drug comparison--cm; polyglactin--pharmaceutics--pr; vitamin D --pharmaceutics--pr; apatite--pharmaceutics--pr; calcium phosphate --pharmaceutics--pr; antibiotic agent--pharmaceutics--pr; unclassified drug

MEDICAL DESCRIPTORS:

\*biodegradable implant

shellfish; biomedical technology assessment; wound dressing; drug delivery system; tissue engineering; Asia; Japan; Korea; Singapore; Taiwan; Thailand ; China; medical research; implant; bone development; rabbit; biocompatibility; bone prosthesis; extracellular matrix; cell regeneration; freeze drying; wound healing; skin; tissue regeneration; polymorphonuclear cell; cell infiltration; tensile strength; mycelium; infection control; burn; ulcer; wound; hydrogel; inflammation; mouth ulcer--drug therapy--dt; drug release; hydrophobicity; drug formulation; nanoparticle; film; liver; particle size; human; nonhuman; article; priority journal

DRUG TERMS (UNCONTROLLED): vinachitin

CAS REGISTRY NO.: 1398-61-4 (chitin); 9012-76-4 (chitosan); 1306-06-5, 51198-94-8 ( hydroxyapatite ); 1305-78-8 (calcium oxide); 13397-26-7, 13701-58-1, 14791-73-2, 471-34-1 (calcium carbonate); 9000-70-8 ( gelatin); 28961-37-7, 29894-36-8, 9005-32-7, 9005-38-3 (alginic acid); 7512-17-6 (n acetylglucosamine); 31799-91-4, 9004-61-9, 9067-32-7 ( hyaluronic acid); 25322-68-3 (macrogol); 22199-08-2 (sulfadiazine silver); 76-25-5 (triamcinolone acetonide); 10344-93-1, 79-10-7 ( acrylic acid); 53570-76-6, 68-19-9, 8064-09-3 (cyanocobalamin); 26780-50-7, 34346-01-5 (polyglactin); 64476-38-6 (apatite); 10103-46-5, 13767-12-9, 14358-97-5, 7758-87-4 (calcium phosphate)

SECTION HEADINGS:

- 027 Biophysics, Bioengineering and Medical Instrumentation
- 033 Orthopedic Surgery
- 037 Drug Literature Index
- 039 Pharmacy

DRUG DESCRIPTORS:

biomaterial--pharmaceutics--pr; hydroxyapatite ; calcium oxide; bone cement; calcium carbonate; gelatin; alginic acid--pharmaceutics--pr; n acetylglucosamine; hyaluronic acid; keratin --pharmaceutics--pr; keratin --pharmacology--pd; macrogol--pharmaceutics--pr; sulfadiazine silver --pharmaceutics--pr; sulfadiazine silver--pharmacology--pd; chitin derivative...

...CAS REGISTRY NO.: 51198-94-8 ( hydroxyapatite ); 1305-78-8 (calcium oxide); 13397-26-7...

3/9,K/25 (Item 3 from file: 73)

DIALOG(R) File 73:EMBASE

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05755932 EMBASE No: 1994169048

Care of the child with tympanostomy tubes: A visual guide for the pediatrician

Isaacson G.; Rosenfeld R.M.  
Dept of Pediatric Otolaryngology, St. Christopher's Hosp. for Children,  
Erie Ave at Front St, Philadelphia, PA 19134-1095 United States  
Pediatrics ( PEDIATRICS ) (United States) 1994, 93/6 I (924-929)  
CODEN: PEDIA ISSN: 0031-4005  
DOCUMENT TYPE: Journal; Article  
LANGUAGE: ENGLISH  
BRAND NAME/MANUFACTURER NAME: cortisporin otic  
DRUG DESCRIPTORS:  
antibiotic agent--drug administration--ad; antibiotic agent--drug therapy  
--dt; beta lactamase; calcium--endogenous compound--ec; chlorine;  
ciprofloxacin--drug therapy--dt; ciprofloxacin--drug administration--ad;  
collagen--endogenous compound--ec; cortisporin--drug therapy--dt;  
cortisporin--drug administration--ad; gentamicin--drug administration--ad;  
gentamicin--drug therapy--dt; hydrocortisone--drug administration--ad;  
hydrocortisone--drug combination--cb; hydrocortisone--drug therapy--dt;  
hydrogen peroxide; **hydroxyapatite**; keratin --endogenous compound--ec;  
metal; neomycin--drug combination--cb; neomycin--drug therapy--dt; neomycin  
--drug administration--ad; phosphate--endogenous compound--ec; plastic;  
polymyxin b--drug therapy--dt; polymyxin b--drug combination--cb; polymyxin  
b--drug administration--ad; tobramycin--drug therapy--dt; tobramycin--drug  
administration--ad  
MEDICAL DESCRIPTORS:  
\*tympanostomy tube  
article; child care; cholesteatoma--complication--co; cholesteatoma  
--etiology--et; cholesteatoma--surgery--su; eardrum; eardrum perforation  
--complication--co; human; microscope; myringotomy; oral drug  
administration; otitis media--drug therapy--dt; otitis media--surgery--su;  
otorrhea--drug therapy--dt; otorrhea--diagnosis--di; otorrhea--complication  
--co; otoscopy; patient referral; pediatrician; perception deafness  
--etiology--et; physical examination; priority journal; reflectometry;  
sclerosis--complication--co; sclerosis--diagnosis--di; topical drug  
administration; tympanometry; tympanoplasty  
CAS REGISTRY NO.: 9073-60-3 (beta lactamase); 7440-70-2 (calcium);  
13981-72-1, 7782-50-5 (chlorine); 85721-33-1 (ciprofloxacin); 9007-34-5

SECTION HEADINGS:

- 004 Microbiology: Bacteriology, Mycology, Parasitology and Virology
- 007 Pediatrics and Pediatric Surgery
- 011 Otorhinolaryngology
- 027 Biophysics, Bioengineering and Medical Instrumentation
- 037 Drug Literature Index

DRUG DESCRIPTORS:

...dt; hydrocortisone--drug administration--ad; hydrocortisone--drug  
combination--cb; hydrocortisone--drug therapy--dt; hydrogen peroxide;  
**hydroxyapatite**; keratin --endogenous compound--ec; metal; neomycin--drug  
combination--cb; neomycin--drug therapy--dt; neomycin--drug administration  
...

...CAS REGISTRY NO.: 51198-94-8 (**hydroxyapatite**); 11004-65-2...

DIALOG(R) File 73:EMBASE  
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01465092 EMBASE No: 1979186084  
**Osteoarthrosis and its treatment**  
Pehlivanov D.  
Res. Inst. Int. Dis. Pharmacol., Med. Acad., Sofia Bulgaria  
MBI Medico-Biologic Information ( MBI MED.-BIOL. INF. ) (Bulgaria) 1979  
NO 2/- (3-7)  
CODEN: MBIFB  
DOCUMENT TYPE: Journal  
LANGUAGE: ENGLISH

DRUG DESCRIPTORS:

\*chondroitin sulfate; \*chondroitin 4 sulfate; \*chondroitin 6 sulfate; \*  
glycine; \*glycosaminoglycan; \*hyaluronic acid; \* hydroxyapatite ; \*  
hydroxyproline; \* keratin ; \*leucine; \*mucin; \*serine; \*threonine; \*  
tyrosine

MEDICAL DESCRIPTORS:

\*arthrosis  
cartilage degeneration; drug therapy; drug administration; intramuscular  
drug administration; therapy; bone; joint

MEDICAL TERMS (UNCONTROLLED): mucarthrin

CAS REGISTRY NO.: 9007-28-7, 9082-07-9 (chondroitin sulfate); 24967-93-9 (chondroitin 4 sulfate); 25322-46-7 (chondroitin 6 sulfate); 56-40-6, 6000-43-7, 6000-44-8 (glycine); 31799-91-4, 9004-61-9, 9067-32-7 (hyaluronic acid); 1306-06-5, 51198-94-8 ( hydroxyapatite ); 51-35-4, 6912-67-0 (hydroxyproline); 61-90-5, 7005-03-0 (leucine); 56-45-1, 6898-95-9 (serine); 36676-50-3, 72-19-5 (threonine); 16870-43-2, 55520-40-6, 60-18-4 (tyrosine)

SECTION HEADINGS:

037 Drug Literature Index

DRUG DESCRIPTORS:

\*chondroitin sulfate; \*chondroitin 4 sulfate; \*chondroitin 6 sulfate; \*  
glycine; \*glycosaminoglycan; \*hyaluronic acid; \* hydroxyapatite ; \*  
hydroxyproline; \* keratin ; \*leucine; \*mucin; \*serine; \*threonine; \*  
tyrosine

...CAS REGISTRY NO.: 51198-94-8 ( hydroxyapatite ); 51-35-4...

3/9,K/27 (Item 1 from file: 94)

DIALOG(R) File 94:JICST-EPlus  
(c)2006 Japan Science and Tech Corp(JST). All rts. reserv.

04402979 JICST ACCESSION NUMBER: 99A0568994 FILE SEGMENT: JICST-E  
**Formation of Hydroxyapatite Layer on Plasma Treated and Grafted Textile Fibers.**  
HIROTSU TOSHIHIRO (1); TSUJISAKA TOSHIHIRO (2); KURAHASHI MASAO (3)  
(1) National Inst. Materials and Chemical Res.; (2) Nara Prefectural Inst.  
Ind. Technol.; (3) Kyoto Munic. Text. Res. Inst.  
Sen'i Gakkai Yokoshu(Sen'i Gakkai Preprints), 1999, VOL.1999, PAGE.G.176,  
REF.2

JOURNAL NUMBER: L1827AAB

UNIVERSAL DECIMAL CLASSIFICATION: 677.027

LANGUAGE: Japanese COUNTRY OF PUBLICATION: Japan

DOCUMENT TYPE: Conference Proceeding

ARTICLE TYPE: Short Communication

MEDIA TYPE: Printed Publication

DESCRIPTORS: hydroxyapatite ; fiber modification; surface treatment; plasma processing; graft copolymerization; cotton(fiber); wool; polyester fiber; nylon fiber; acrylic fiber; silk; vinyl compound; aliphatic carboxylic acid; unsaturated carboxylic acid

BROADER DESCRIPTORS: apatite; phosphate mineral; mineral(geology); reforming; treatment; copolymerization; polymerization; chemical reaction; seed hair fiber; vegetable fiber; cellulosic fiber; fiber; natural fiber; keratin fiber; animal fiber; protein fiber; synthetic fiber; man-made fiber; polyamide fiber; olefin compound; carboxylic acid

CLASSIFICATION CODE(S) : YM02050Z

**Formation of Hydroxyapatite Layer on Plasma Treated and Grafted Textile Fibers.**

DESCRIPTORS: hydroxyapatite ;  
...BROADER DESCRIPTORS: keratin fiber

3/9,K/28 (Item 2 from file: 94)

DIALOG(R) File 94:JICST-EPlus

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02666535 JICST ACCESSION NUMBER: 96A0269596 FILE SEGMENT: JICST-E  
**The Mineralization of Crystalline Inorganic Components in Japanese Serow Horn.**

HASHIGUCHI K (1); HASHIMOTO K (1)

(1) Hamamatsu Univ. School of Medicine, Shizuoka, JPN

Okajimas Folia Anat Jpn, 1995, VOL.72,NO.5, PAGE.235-243, FIG.5, TBL.1,  
REF.22

JOURNAL NUMBER: F0730AAJ ISSN NO: 0030-154X CODEN: OFAJA

UNIVERSAL DECIMAL CLASSIFICATION: 591.177.05+591.471

LANGUAGE: English COUNTRY OF PUBLICATION: Japan

DOCUMENT TYPE: Journal

ARTICLE TYPE: Original paper

MEDIA TYPE: Printed Publication

ABSTRACT: The Japanese serow (*Capricornis crispus*) is protected as a special natural monument in Japan. The ring count of the soft X-ray photographs of Japanese serow horn was found to be a useful criteria to determine the ages exactly. The mineralization process in Japanese serow horn was examined microscopic, ICP and X-ray diffraction methods. The incremental lines appeared as light and dark layers in the section stained for fuchsin and methylen blue. Mineral depositions were observed among the keratin fibers, no matrix vesicle in the electron dense regions. X-ray diffraction pattern of crystalline inorganic components in Japanese serow horn was determined as .BETA.-tricalcium phosphate (TCP), hydroxyapatite (HA) and unknown phase. ICP measurement was also carried out. The horn contained trace elements of K besides Na, Ca, Fe and P. The Ca/P molar was found to be 2.9. The ratio was much higher than the theoretical value of HA. Presumably, keratin was the seed which might be related to mineralization and higher Ca activity was detected in the initial phase of epitaxial growth. Analytical results of the measurement of trace elements in Japanese serow horn by using ICP method seemed to be correlated with the evaluation of environmental conditions. The present study indicated that the mineralization of Japanese serow horn directly related with deposition Ca-deficient HA among the keratin fibers. (author abst.)

DESCRIPTORS: Capricornis crispus; horn(animal tissue); calcification(physiology); keratin fiber; hydroxyapatite ; calcium; phosphorus; minor component; X-ray diffraction; ICP(analys); electron microscopy; optical microscopy; mineral metabolism

BROADER DESCRIPTORS: Bovidae; Ruminantia; Artiodactyla; Mammalia; Vertebrata; animal; head(body region); body region; variation; animal fiber; protein fiber; fiber; natural fiber; apatite; phosphate mineral; mineral(geology); alkaline earth metal; metallic element; element; fourth row element; third row element; nitrogen group element; component; X-ray scattering; electromagnetic wave scattering; scattering; diffraction; coherent scattering; plasma spectrochemical analysis; instrumental analysis; analysis(separation); analysis; microscopy; observation and view; metabolism

CLASSIFICATION CODE(S): EJ10030R

...ABSTRACT: in the section stained for fuchsin and methylen blue. Mineral depositions were observed among the keratin fibers, no matrix vesicle in the electron dense regions. X-ray diffraction pattern of crystalline inorganic components in Japanese serow horn was determined as .BETA.-tricalcium phosphate (TCP), hydroxyapatite (HA) and unknown phase. ICP measurement was also carried out. The horn contained trace elements...

...be 2.9. The ratio was much higher than the theoretical value of HA. Presumably, keratin was the seed which might be related to mineralization and higher Ca activity was detected...

...the mineralization of Japanese serow horn directly related with deposition Ca-deficient HA among the keratin fibers. (author abst.)

...DESCRIPTORS: keratin fiber...

... hydroxyapatite ;

3/9,K/29 (Item 3 from file: 94)

DIALOG(R) File 94:JICST-EPlus

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01515644 JICST ACCESSION NUMBER: 92A0133982 FILE SEGMENT: JICST-E  
Cell biological studies on biocompatibility of hydroxyapatite:An analysis of cellular activities expressed by established human gingival cells cultured on the hydroxyapatite .

ISHIKAWA KEIKO (1)

(1) Okazaki National Res. Inst.

Shika Kiso Igakkai Zasshi(Japanese Journal of Oral Biology), 1991,  
VOL.33,NO.6, PAGE.513-533, FIG.17, REF.36

JOURNAL NUMBER: Y0018AAZ ISSN NO: 0385-0137

UNIVERSAL DECIMAL CLASSIFICATION: 616.314-7

LANGUAGE: Japanese COUNTRY OF PUBLICATION: Japan

DOCUMENT TYPE: Journal

ARTICLE TYPE: Original paper

MEDIA TYPE: Printed Publication

ABSTRACT: Although hydroxyapatite has been widely applied in the field of dental medicine, there is the argument about clinical complaints due to infection and inflammation after its implantation. In the present study I have attempted to establish a useful in vitro system to evaluate the biocompatibility of hydroxyapatite and other biomaterials. I have firstly established a human gingival fibroblast line designated as

HGF-22 by clonal selection techniques and two human gingival epithelial cell lines designated as HGE-15vI and HGE-15vII, respectively, by transfection of SV40-T genes. All these established cells grow actively and maintain their cellular characteristics with stability. HGE-15vI cells conserve much higher activity to synthesize keratin molecules than HGE-15vII cells. Using HGF-22 and HGE-15vI cells I have quantitatively analyzed their activities represented by cell adhesion, spreading, growth and differentiation when cultured on the surface of a hydroxyapatite disc, which is thin enough to observe their living conditions by a phase contrast microscope, and on the plastic surface of commercial cell culture dishes. From results obtained it is possible to conclude that the established cell lines can provide highly useful experimental tools for basic studies of clinically used biomaterials such as hydroxyapatite. (author abst.)

DESCRIPTORS: histocompatibility; hydroxyapatite; gingiva; fibroblast; epithelium; human(primates); cultured cell; cell proliferation; differentiation antigen; electron microscopy; transfection; SV40 virus; adhesion(surface chemistry)

BROADER DESCRIPTORS: transplantation immunity; immunological reaction; reaction; biocompatibility; property; apatite; phosphate mineral; mineral(geology); periodontium; oral cavity; digestive organ; blast cell; cell(cytology); epithelial tissue; animal tissue; biomedical tissue; organization; histomembrane; membrane and film; cell physiology; multiplication(biology); surface antigen; antigen; microscopy; observation and view; gene introduction; gene manipulation; genetic technique; technology; operation(processsing); Polyomavirus; Papovaviridae; DNA virus; virus; microorganism; tumor virus; animal virus

CLASSIFICATION CODE(S): GT06000B

...hydroxyaptite:An analysis of cellular activities expressed by established human gingival cells cultured on the hydroxyapatite .

ABSTRACT: Although hydroxyapatite has been widely applied in the field of dental medicine, there is the argument about...

...I have attempted to establish a useful in vitro system to evaluate the biocompatibility of hydroxyapatite and other biomaterials. I have firstly established a human gingival fibroblast line designated as HGF

... maintain their cellular characteristics with stability. HGE-15vI cells conserve much higher activity to synthesize keratin molecules than HGE-15vII cells. Using HGF-22 and HGE-15vI cells I have quantitatively

...represented by cell adhesion, spreading, growth and differentiation when cultured on the surface of a hydroxyapatite disc, which is thin enough to observe their living conditions by a phase contrast microscope...

...can provide highly useful experimental tools for basic studies of clinically used biomaterials such as hydroxyapatite . (author abst.)

...DESCRIPTORS: hydroxyapatite ;

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0000155296 IP ACCESSION NO: 5977975

Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation

Tachibana, A; Kaneko, S; Tanabe, T; Yamauchi, K

Department of Applied and Bioapplied Chemistry, Graduate School of Engineering, Osaka City University, Sugimoto 3-3-138, Sumiyoshi-ku, Osaka 558- 8585, Japan, [mailto:[tatibana@bioa.eng.osaka-cu.ac.jp](mailto:tatibana@bioa.eng.osaka-cu.ac.jp)]

Biomaterials, v 26, n 3, p 297-302, January 2005

PUBLICATION DATE: 2005

PUBLISHER: Elsevier Science Ltd., The Boulevard Langford Lane Kidlington Oxford OX5 1GB UK, [mailto:[nlinfo-f@elsevier.nl](mailto:nlinfo-f@elsevier.nl)], [URL:<http://www.elsevier.nl>]

DOCUMENT TYPE: Journal Article

RECORD TYPE: Abstract

LANGUAGE: English

SUMMARY LANGUAGE: English

ISSN: 0142-9612

DOI: 10.1016/j.biomaterials.2004.02.032

FILE SEGMENT: BioEngineering Abstracts

ABSTRACT:

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium phosphate hybrid biomaterials were described. Firstly, the CaP-precipitated sponges were obtained by only the immersion of the carboxyl-sponges, chemically introduced with high amount of carboxyl groups on the sponges, in calcium and phosphate ions containing buffers such as PBS(+) for only 1-3 days. Neither sponge, introduced with amino or amido groups or non-treated, gave significant calcium phosphate precipitation. The carboxyl-sponges were mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast differentiation marker, suggested that both hybrid sponges, CaP-precipitated and trapped sponges, alter the differentiation pattern of preosteoblasts, MC3T3-E1.

DESCRIPTORS: Calcium phosphate; Hybrids; Keratin ; Osteoblastogenesis; Hydroxyapatite ; Biomaterials; Osteoblasts; Precipitation;

Calcification; Wool; Immersion; Alkaline phosphatase

SUBJ CATG: 110, Biomedical Materials & Tissue Engineering

Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation

ABSTRACT:

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The

hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium...

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DESCRIPTORS: Calcium phosphate; Hybrids; Keratin ; Osteoblastogenesis; Hydroxyapatite ; Biomaterials; Osteoblasts; Precipitation; Calcification; Wool; Immersion; Alkaline phosphatase

3/9,K/31 (Item 1 from file: 144)

DIALOG(R) File 144:Pascal

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16831162 PASCAL No.: 04-0490140

The mechanical efficiency of natural materials

WEGST U G K; ASHBY M. F

Max-Planck-Institut fuer Metallforschung, Heisenbergstrasse 3, 70569, Stuttgart, Germany; Engineering Design Centre, Engineering Department, University of Cambridge, Trumpington Street, Cambridge CB2 1PZ, United Kingdom

Journal: Philosophical magazine : (2003. Print), 2004, 84 (21) 2167-2181

ISSN: 1478-6435 Availability: INIST-134A3; 354000120031670030

No. of Refs.: 10 ref.

Document Type: P (Serial) ; A (Analytic)

Country of Publication: United Kingdom

Language: English

The materials of nature, for example cellulose, lignin, keratin, chitin, collagen and hydroxyapatite , and the structures made from them, for example bamboo, wood, antler and bone, have a remarkable range of mechanical properties. These can be compared by presenting them as material property charts, well known for the materials of engineering. Material indices (significant combinations of properties) can be plotted on to the charts, identifying materials with extreme values of an index, suggesting that they have evolved to carry particular modes of loading, or to sustain large tensile or flexural deformations, without failure. This paper describes a major revision and update of a set of property charts for natural material published some 8 years ago by Ashby et al. with examples of their use to study mechanical efficiency in nature.

English Descriptors: Reviews; Deformation; Microstructure; Young modulus; Mechanical strength; Fracture toughness; Cellulose; Lignin; Keratin ; Chitin; Collagen; Hydroxyapatite ; Wood

Broad Descriptors: Organic compounds; Compose organique

French Descriptors: Article synthese; Deformation; Microstructure; Module Young; Resistance mecanique; Tenacite; Cellulose; Lignine; Keratine; Chitine; Collagene; Apatite hydroxylee; Bois; 6220F

Classification Codes: 001B60B20F

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The materials of nature, for example cellulose, lignin, keratin, chitin, collagen and hydroxyapatite, and the structures made from them, for example bamboo, wood, antler and bone, have a...

English Descriptors: Reviews; Deformation; Microstructure; Young modulus; Mechanical strength; Fracture toughness; Cellulose; Lignin; Keratin; Chitin; Collagen; Hydroxyapatite; Wood

3/9,K/32 (Item 2 from file: 144)

DIALOG(R) File 144:Pascal  
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15898483 PASCAL No.: 03-0037198  
**Isolation of Thermoanaerobacter keratinophilus sp. nov., a novel thermophilic, anaerobic bacterium with keratinolytic activity**  
RIESSEN Sabine; ANTRANIKIAN Garabed  
Institute of Technical Microbiology, Technical University  
Hamburg-Harburg, Kasernenstr. 12, 21073 Hamburg, Germany  
Journal: Extremophiles, 2001, 5 (6) 399-408  
ISSN: 1431-0651 Availability: INIST-26559; 354000103519230050  
No. of Refs.: 1 p.1/4  
Document Type: P (Serial); A (Analytic)  
Country of Publication: Japan  
Language: English  
Several thermophilic anaerobic bacteria with keratinolytic activity growing at temperatures between 50 Degree C and 90 Degree C were isolated from samples collected on the island of Sao Miguel in the Azores (Portugal). On the basis of morphological, physiological, and 16S rDNA studies, the isolate 2KXI was identified as a new species of the genus Thermoanaerobacter, designated Thermoanaerobacter keratinophilus. This strain, which grows optimally at 70 Degree C, pH 7.0, and 0.5% NaCl, is the first member of the genus Thermoanaerobacter that has been described for its ability to degrade native keratin. Around 70% of native wool was solubilized after 10 days of incubation under anaerobic conditions. The strain was shown to possess intracellular and extracellular proteases optimally active at 60 Degree C, pH 7.0, and 85 Degree C, pH 8.0, respectively. Keratin hydrolysis was demonstrated in vitro using a sodium dodecyl sulfate gel containing feather meal. The extracellular protease responsible for breaking down keratin fibers was purified to homogeneity in only one step by applying hydroxyapatite column chromatography. The enzyme belongs to the serine-type proteases and has a molecular mass of 135 kDa.

English Descriptors: Thermoanaerobacter; Thermophily; Anaerobe; Temperature; Island; Azores; Portugal; Ribosomal DNA; New genus; pH; Isolate; New species  
Broad Descriptors: Bacteria; Atlantic Ocean Islands; Europe; Biotope; Bacterie; Iles Atlantiques; Europe; Biotope; Bacteria; Islas Atlantico; Europa; Biotopo

French Descriptors: Thermoanaerobacter; Thermophilie; Anaerobie; Temperature; Ile; Acores; Portugal; DNA ribosomique; Genre nouveau; pH; Isolat; Espece nouvelle

Classification Codes: 002A05B02; 002A05B09

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...member of the genus *Thermoanaerobacter* that has been described for its ability to degrade native keratin. Around 70% of native wool was solubilized after 10 days of incubation under anaerobic conditions...

...at 60 Degree C, pH 7.0, and 85 Degree C, pH 8.0, respectively. Keratin hydrolysis was demonstrated in vitro using a sodium dodecyl sulfate gel containing feather meal. The extracellular protease responsible for breaking down keratin fibers was purified to homogeneity in only one step by applying hydroxyapatite column chromatography. The enzyme belongs to the serine-type proteases and has a molecular mass...

? t s3/9,k/33-40

3/9,K/33 (Item 1 from file: 155)  
DIALOG(R) File 155: MEDLINE(R)  
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21634580 PMID: 16823809

Structure of white rhinoceros (*Ceratotherium simum*) horn investigated by X-ray computed tomography and histology with implications for growth and external form.

Hieronymus Tobin L; Witmer Lawrence M; Ridgely Ryan C  
Department of Biological Sciences, Ohio University, Athens, Ohio 45701,  
USA. Th108702@ohio.edu

Journal of morphology (United States) Oct 2006, 267 (10) p1172-6,  
ISSN 0362-2525--Print Journal Code: 0406125

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: In Process

Subfile: INDEX MEDICUS

The nasal and frontal horns of two individuals of *Ceratotherium simum* were examined by x-ray computed tomography (CT scanning), gross observation of sectioned horn, and light microscopy of histological sections of the horn tissue. CT scans of both sets of horns reveal a periodic banding pattern that is evident upon gross observation of sections as darker bands of tissue. The overlap of these bands in both histological and CT slices suggests the presence of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely hydroxyapatite or octocalcium phosphate). The distribution of these two components in the horns is hypothesized to contribute to the differential wear patterns that produce the characteristic sweeping conical shape of rhinoceros horn from what otherwise (in the absence of wear and UV exposure) would be cylindrical blocks of constantly growing cornified papillary epidermis. Although extant rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of keratin -and-bone horns such as those of bovid artiodactyls, the tissue structures that make up the horn are strikingly convergent with other examples of papillary cornified epidermis found in horses, artiodactyls, cetaceans, and birds. Copyright (c) 2006 Wiley-Liss, Inc.

Record Date Created: 20060904

... of both a photoabsorbent component (melanin) and a radiodense component (calcium phosphate salts, most likely hydroxyapatite or octocalcium phosphate). The distribution of these two components in the horns is hypothesized to...

... rhinocerotids are unique in possessing a massive entirely keratinous horn that approximates the functions of keratin -and-bone horns such as those of bovid artiodactyls, the tissue structures that make up...

3/9,K/34 (Item 2 from file: 155)  
DIALOG(R) File 155: MEDLINE(R)  
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19707234 PMID: 16211607

Applications of X-ray powder diffraction in materials chemistry.

Skakle Jan

Department of Chemistry, University of Aberdeen, Meston Walk, Aberdeen AB24 3UE, Scotland, United Kingdom. j.skakle@abdn.ac.uk

Chemical record (New York, N.Y.) (United States) 2005, 5 (5) p252-62, ISSN 1527-8999--Print Journal Code: 101085550

Publishing Model Print

Document type: Journal Article; Review

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subfile: INDEX MEDICUS

X-ray powder diffraction is a standard technique in materials chemistry, yet it is often still used in the laboratory as a "one-hit" technique, e.g. for fingerprinting and following the progress of reactions. It is important, however, that the wealth of information available from powder data is not overlooked. While it is only possible here to scratch the surface of possibilities, a range of examples from our research is used to emphasize some of the more accessible techniques and to highlight successes as well as potential problems. The first example is the study of solid solution formation in the oxide systems Ba(3-3x)La(2x)V2O8 and Sr(4-x)Ba(x)Mn3O10 and in the silicate- hydroxyapatite bioceramic, Ca10(PO4)6-x(SiO4)x(OH)2-x. Database mining is also explored, using three phases within the pseudobinary phase diagram Li3SbO4-CuO as examples. All three phases presented different challenges: the structure of Li3SbO4 had been previously reported in higher symmetry than was actually the case, Li3Cu2SbO6 was found to be isostructural with Li2TiO3 but the cation ordering had to be rationalized, and Li3CuSbO5 was believed to be triclinic, presenting challenges in indexing the powder pattern. Quantitative phase analysis is briefly discussed, with the emphasis both on success (determination of amorphous phase content in a novel cadmium arsenate phase) and on possible failure (compositional analysis in bone mineral); the reasons for the problems in the latter are also explored. Finally, the use of an area detector system has been shown to be of value in the study of orientational effects (or lack of them) in non- and partially-ordered biomaterials, including p-HEMA, annulus fibrosis of lumbar discs, and keratin in the horn of cow's hooves. Copyright 2005 The Japan Chemical Journal Forum and Wiley Periodicals, Inc (62 Refs.)

Descriptors: \*Biocompatible Materials--chemistry--CH; \*Powders--chemistry --CH; \*X-Ray Diffraction--methods--MT; Animals; Cattle; Databases; Durapatite--chemistry--CH; Keratin --chemistry--CH; Models, Chemical; Research Support, Non-U.S. Gov't

CAS Registry No.: 0 (Biocompatible Materials); 0 (Powders); 1306-06-5  
(Durapatite); 68238-35-7 (Keratin)  
Record Date Created: 20051013  
Record Date Completed: 20060111

... $3x$ )La( $2x$ )V<sub>2</sub>O<sub>8</sub> and Sr( $4-x$ )Ba( $x$ )Mn<sub>3</sub>O<sub>10</sub> and in the silicate-hydroxyapatite bioceramic, Ca<sub>10</sub>(PO<sub>4</sub>)<sub>6-x</sub>(SiO<sub>4</sub>)<sub>x</sub>(OH)<sub>2-x</sub>. Database mining is also explored...

... in non- and partially-ordered biomaterials, including p-HEMA, annulus fibrosis of lumbar discs, and keratin in the horn of cow's hooves. Copyright 2005 The Japan Chemical Journal Forum and...

; Animals; Cattle; Databases; Durapatite--chemistry--CH; Keratin--chemistry--CH; Models, Chemical; Research Support, Non-U.S. Gov't Chemical Name: Biocompatible Materials; Powders; Durapatite; Keratin

3/9,K/35 (Item 3 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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15250029 PMID: 15262471

Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation.

Tachibana Akira; Kaneko Sumika; Tanabe Toshizumi; Yamauchi Kiyoshi  
Department of Applied and Bioapplied Chemistry, Graduate School of Engineering, Osaka City University, Sugimoto 3-3-138, Sumiyoshi-ku, Osaka 558-8585, Japan. tatibana@biao.eng.osaka-cu.ac.jp

Biomaterials (England) Jan 2005, 26 (3) p297-302, ISSN 0142-9612--  
Print Journal Code: 8100316

Publishing Model Print

Document type: Evaluation Studies; Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subfile: INDEX MEDICUS

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium phosphate hybrid biomaterials were described. Firstly, the CaP-precipitated sponges were obtained by only the immersion of the carboxyl-sponges, chemically introduced with high amount of carboxyl groups on the sponges, in calcium and phosphate ions containing buffers such as PBS(+) for only 1-3 days. Neither sponge, introduced with amino or amido groups or non-treated, gave significant calcium phosphate precipitation. The carboxyl-sponges were mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast differentiation marker, suggested that both hybrid sponges, CaP-precipitated and trapped sponges, alter the differentiation pattern of preosteoblasts, MC3T3-E1.

Descriptors: \*Bone Substitutes--chemistry--CH; \*Durapatite--chemistry--CH; \*Keratin --chemistry--CH; \*Osteoblasts--cytology--CY; \*Osteoblasts--physiology--PH; \*Osteogenesis--physiology--PH; \*Tissue Engineering

--methods--MT; Animals; Biocompatible Materials--chemistry--CH; Cell Differentiation--physiology--PH; Comparative Study; Materials Testing; Mice ; Research Support, Non-U.S. Gov't  
· CAS Registry No.: 0 (Biocompatible Materials); 0 (Bone Substitutes); 1306-06-5 (Durapatite); 68238-35-7 (Keratin)  
Record Date Created: 20040720  
Record Date Completed: 20050215

Rapid fabrication of keratin - hydroxyapatite hybrid sponges toward osteoblast cultivation and differentiation.

Wool keratin sponges were reported to be useful scaffolds for long-term and high-density cell cultivation (J. Biotechnol. 93 (2002) 165). The hybrid of the keratin sponges with calcium phosphate materials gave the additional function. Two rapid fabrication methods for calcium...

... mimics of matrix gamma-carboxyglutamic acid protein, which is responsible for osteoblast calcification. Secondly, the hydroxyapatite particle suspension was added onto carboxyl-sponges to fabricate trapped sponge. The trapped hydroxyapatite particles might interact with keratin protein of the sponge walls. Preliminary experiments measuring the expression of alkaline phosphatase, early osteoblast...

Descriptors: \*Bone Substitutes--chemistry--CH; \*Durapatite--chemistry--CH ; \* Keratin --chemistry--CH; \*Osteoblasts--cytology--CY; \*Osteoblasts --physiology--PH; \*Osteogenesis--physiology--PH; \*Tissue Engineering --methods--MT

Chemical Name: Biocompatible Materials; Bone Substitutes; Durapatite; Keratin

3/9,K/36 (Item 4 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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13519666 PMID: 11778841

Isolation of Thermoanaerobacter keratinophilus sp. nov., a novel thermophilic, anaerobic bacterium with keratinolytic activity.

Riessen S; Antranikian G  
Institute of Technical Microbiology, Technical University  
Hamburg-Harburg, Germany.

Extremophiles - life under extreme conditions (Germany) Dec 2001, 5  
(6) p399-408, ISSN 1431-0651--Print Journal Code: 9706854

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subfile: INDEX MEDICUS; SPACE LIFE SCIENCES

Several thermophilic anaerobic bacteria with keratinolytic activity growing at temperatures between 50 degrees C and 90 degrees C were isolated from samples collected on the island of Sao Miguel in the Azores (Portugal). On the basis of morphological, physiological, and 16S rDNA studies, the isolate 2KXI was identified as a new species of the genus Thermoanaerobacter, designated Thermoanaerobacter keratinophilus. This strain, which grows optimally at 70 degrees C, pH 7.0, and 0.5% NaCl, is the first member of the genus Thermoanaerobacter that has been described for its ability to degrade native keratin. Around 70% of native wool was solubilized after 10 days of incubation under anaerobic conditions. The strain was shown to possess intracellular and extracellular proteases

optimally active at 60 degrees C, pH 7.0, and 85 degrees C, pH 8.0, respectively. **Keratin** hydrolysis was demonstrated in vitro using a sodium dodecyl sulfate gel containing feather meal. The extracellular protease responsible for breaking down **keratin** fibers was purified to homogeneity in only one step by applying **hydroxyapatite** column chromatography. The enzyme belongs to the serine-type proteases and has a molecular mass of 135 kDa.

Descriptors: \*Bacillaceae--isolation and purification--IP; \*Bacillaceae--metabolism--ME; \*Bacteria, Anaerobic--isolation and purification--IP; \*Bacteria, Anaerobic--metabolism--ME; \*Keratin --metabolism--ME; Bacillaceae--classification--CL; Bacillaceae--genetics--GE; Bacteria, Anaerobic--classification--CL; Bacteria, Anaerobic--genetics--GE; Biodegradation; DNA, Bacterial--genetics--GE; DNA, Ribosomal--genetics--GE; Heat; Peptide Hydrolases--chemistry--CH; Peptide Hydrolases--metabolism--ME; Phylogeny; Research Support, Non-U.S. Gov't; Textiles

CAS Registry No.: 0 (DNA, Bacterial); 0 (DNA, Ribosomal); 68238-35-7 (Keratin)

Enzyme No.: EC 3.4.- (Peptide Hydrolases); EC 3.4.- (keratinase)

Record Date Created: 20020107

Record Date Completed: 20020709

...member of the genus *Thermoanaerobacter* that has been described for its ability to degrade native **keratin**. Around 70% of native wool was solubilized after 10 days of incubation under anaerobic conditions...

...at 60 degrees C, pH 7.0, and 85 degrees C, pH 8.0, respectively. **Keratin** hydrolysis was demonstrated in vitro using a sodium dodecyl sulfate gel containing feather meal. The extracellular protease responsible for breaking down **keratin** fibers was purified to homogeneity in only one step by applying **hydroxyapatite** column chromatography. The enzyme belongs to the serine-type proteases and has a molecular mass...

...Descriptors: purification--IP; \*Bacillaceae--metabolism--ME; \*Bacteria, Anaerobic--isolation and purification--IP; \*Bacteria, Anaerobic--metabolism--ME; \*Keratin --metabolism--ME

Chemical Name: DNA, Bacterial; DNA, Ribosomal; Keratin ; Peptide Hydrolases; keratinase

3/9,K/37 (Item 5 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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11650877 PMID: 11063033

Identification of tuftelin- and amelogenin-interacting proteins using the yeast two-hybrid system.

Paine C T; Paine M L; Snead M L  
University of Southern California, School of Dentistry, Center for Craniofacial Molecular Biology, Los Angeles 90033, USA.

Connective tissue research (ENGLAND) 1998, 38 (1-4)  
p257-67; discussion 295-303, ISSN 0300-8207--Print Journal Code: 0365263  
Contract/Grant No.: DE 06988; DE; NIDCR; DE 07211; DE; NIDCR; DE 11704;  
DE; NIDCR

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subfile: INDEX MEDICUS

Biomineralization of enamel is a complex process that involves the eventual replacement of an extracellular protein matrix by hydroxyapatite crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin and ameloblastin. Assembly of the enamel extracellular matrix from these component proteins is believed to be critical in producing a matrix competent to undergo mineral replacement. Enamel formation is a complex process and additional proteins are likely to have a role in the assembly of the extracellular matrix. In order to identify additional proteins involved in the assembly process, the yeast two-hybrid system developed by Fields and Song (1989) has been implemented. This system allows for the identification of unknown proteins that interact with proteins of interest. Typically a known protein is used as "bait" to screen a cDNA expression library of interest. In our studies, tuftelin or amelogenin have been used to screen a mouse tooth library produced from one day old pups. A library screening of six million clones with amelogenin as bait resulted in eleven positive clones all of which show high homology to the human leukocyte antigen-B (HLA-B) associated transcript (BAT) family of genes. A library screening of one million clones using tuftelin as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either keratin K5 or keratin K6, four are constitutively expressed and the remaining seven are novel. Further characterization of the proteins shown to interact with amelogenin or tuftelin may shed additional light on this complex process of enamel matrix assembly.

Descriptors: \*Dental Enamel Proteins--metabolism--ME; Animals; Dental Enamel Proteins--genetics--GE; HLA-B Antigens--metabolism--ME; Humans; Mice ; Research Support, U.S. Gov't, P.H.S.; Two-Hybrid System Techniques; Yeasts

CAS Registry No.: 0 (Dental Enamel Proteins); 0 (HLA-B Antigens); 0 (TUFT1 protein, human); 0 (amelogenins)

Record Date Created: 20001130

Record Date Completed: 20001130

... is a complex process that involves the eventual replacement of an extracellular protein matrix by hydroxyapatite crystallites. To date four different enamel matrix proteins have been identified; the amelogenins, tuftelin, enamelin...

... as the bait identified twenty-one tuftelin-interacting proteins. Ten of these proteins are either keratin K5 or keratin K6, four are constitutively expressed and the remaining seven are novel. Further characterization of the...

3/9,K/38 (Item 6 from file: 155)  
DIALOG(R) File 155: MEDLINE(R)  
(c) format only 2006 Dialog. All rts. reserv.

11555392 PMID: 9395922

Cytokeratin 8 and 19 as antigens recognized by adenocarcinoma-reactive human monoclonal antibody AE6F4.

Ichikawa A; Tachibana H; Kawamoto S; Kamei M; Honjoh T; Hashizume S; Shirahata S

Graduate School of Genetic Resources Technology, Faculty of Agriculture, Kyushu University, Fukuoka, Japan.

Human antibodies (UNITED STATES) 1997, 8 (4) p195-202, ISSN

1093-2607--Print Journal Code: 9711270

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subfile: INDEX MEDICUS

The human monoclonal antibody (MAb) AE6F4 is secreted by a human-human hybridoma line established from the in vitro immunization of normal human peripheral blood lymphocytes with the human lung adenocarcinoma cell line, A549. This MAb is strongly reactive to lung cancer tissues. In the previous study, the antigens recognized by the MAb AE6F4 were purified from A549 cells and identified as 14-3-3 protein and 31 kDa cytosolic phospholipase A2 (cPLA2). The MAb AE6F4 also binds two kinds of antigens (53 kDa and 40 kDa), which are not related to 14-3-3 protein or 31 kDa cPLA2, in the human breast adenocarcinoma cell line, MCF-7. We purified a 38 kDa antigen, which is a degradation product of 53 kDa antigen from breast adenocarcinoma MCF-7 cells using ion-exchange and hydroxyapatite column chromatography. Two partial amino acid sequences of the purified 38 kDa antigen showed 95-100% homology to human cytokeratin 8 (CK8). Two-dimensional gel electrophoresis and immunoblot analysis of intermediate filament fraction separated from MCF-7 cells demonstrated that the 53 kDa and 40 kDa antigens were CK8 and CK19, respectively. Antigenic determinants on CK8 and CK19 recognized by the MAb AE6F4 were resistant to sodium periodate treatment, although antigenic determinant on 31 kDa antigen (14-3-3 protein and(or) cPLA2) was sensitive to this treatment. These results suggest that the MAb AE6F4 reacts with both carbohydrate and peptide antigenic determinants.

Descriptors: \*Adenocarcinoma--immunology--IM; \*Antibodies, Monoclonal; \*Antigens; \* Keratin --immunology--IM; Adenocarcinoma--diagnosis--DI; Amino Acid Sequence; Antibodies, Monoclonal--diagnostic use--DU; Antigens --chemistry--CH; Antigens--genetics--GE; Carbohydrates--chemistry--CH; Carbohydrates--immunology--IM; Epitopes--chemistry--CH; Epitopes --genetics--GE; Humans; Hybridomas--immunology--IM; Keratin --chemistry --CH; Keratin --genetics--GE; Molecular Sequence Data; Molecular Weight; Peptides--chemistry--CH; Peptides--genetics--GE; Peptides--immunology--IM ; Tumor Cells, Cultured

CAS Registry No.: 0 (Antibodies, Monoclonal); 0 (Antigens); 0 (Carbohydrates); 0 (Epitopes); 0 (Peptides); 68238-35-7 (Keratin)

Record Date Created: 19980115

Record Date Completed: 19980115

...product of 53 kDa antigen from breast adenocarcinoma MCF-7 cells using ion-exchange and hydroxyapatite column chromatography. Two partial amino acid sequences of the purified 38 kDa antigen showed 95...

Descriptors: \*Adenocarcinoma--immunology--IM; \*Antibodies, Monoclonal; \*Antigens; \* Keratin --immunology--IM...; chemistry--CH; Carbohydrates --immunology--IM; Epitopes--chemistry--CH; Epitopes--genetics--GE; Humans; Hybridomas--immunology--IM; Keratin --chemistry--CH; Keratin --genetics --GE; Molecular Sequence Data; Molecular Weight; Peptides--chemistry--CH; Peptides--genetics--GE; Peptides--immunology...

Chemical Name: Antibodies, Monoclonal; Antigens; Carbohydrates; Epitopes; Peptides; Keratin

3/9,K/39 (Item 7 from file: 155)

DIALOG(R) File 155: MEDLINE(R)

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10775934 PMID: 8868213

The mineralization of crystalline inorganic components in Japanese serow horn.

Hashiguchi K; Hashimoto K  
Department of Oral Surgery, Hamamatsu University School of Medicine,  
Shizuoka, Japan.

Okajimas folia anatomica Japonica (JAPAN) Dec 1995, 72 (5) p235-43,  
ISSN 0030-154X--Print Journal Code: 0401014

Publishing Model Print

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM

Record type: MEDLINE; Completed

Subfile: INDEX MEDICUS

The Japanese serow (*Capricornis crispus*) is protected as a special natural monument in Japan. The ring count of the soft X-ray photographs of Japanese serow horn was found to be a useful criteria to determine the ages exactly. The mineralization process in Japanese serow horn was examined microscopic, ICP and X-ray diffraction methods. The incremental lines appeared as light and dark layers in the section stained for fuchsin and methylen blue. Mineral depositions were observed among the keratin fibers, no matrix vesicle in the electron dense regions. X-ray diffraction pattern of crystalline inorganic components in Japanese serow horn was determined as beta-tricalcium phosphate (TCP), hydroxyapatite (HA) and unknown phase. ICP measurement was also carried out. The horn contained trace elements of K besides Na, Ca, Fe and P. The Ca/P molar was found to be 2.9. The ratio was much higher than the theoretical value of HA. Presumably, keratin was the seed which might be related to mineralization and higher Ca activity was detected in the initial phase of epitaxial growth. Analytical results of the measurement of trace elements in Japanese serow horn by using ICP method seemed to be correlated with the evaluation of environmental conditions. The present study indicated that the mineralization of Japanese serow horn directly related with deposition Ca-deficient HA among the keratin fibers.

Descriptors: \*Antelopes--metabolism--ME; \*Calcification, Physiologic; \*Horns--metabolism--ME; \*Minerals--metabolism--ME; Animals; Crystallization; Hydroxyapatites--metabolism--ME

CAS Registry No.: 0 (Hydroxyapatites); 0 (Minerals)

Record Date Created: 19961126

Record Date Completed: 19961126

... in the section stained for fuchsin and methylen blue. Mineral depositions were observed among the keratin fibers, no matrix vesicle in the electron dense regions. X-ray diffraction pattern of crystalline inorganic components in Japanese serow horn was determined as beta-tricalcium phosphate (TCP), hydroxyapatite (HA) and unknown phase. ICP measurement was also carried out. The horn contained trace elements...

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... the mineralization of Japanese serow horn directly related with deposition Ca-deficient HA among the keratin fibers.

3/9,K/40 (Item 1 from file: 357)  
DIALOG(R) File 357:Derwent Biotech Res.  
(c) 2007 The Thomson Corp. All rts. reserv.

0375660 DBR Accession No.: 2005-21366 PATENT  
Novel protease having high degradation activity with respect to skin desmosome, in comparison with skin keratin, useful for preparing a skin cleaning agent composition - isolation and purification of a protease from *Bacillus* useful for the preparation of a cosmetic composition for desmosome degradation

AUTHOR: SUZUKI N; UENO J; KIGAWA H

PATENT ASSIGNEE: LION CORP 2005

PATENT NUMBER: JP 2005192403 PATENT DATE: 20050721 WPI ACCESSION NO.: 2005-501991 (200551)

PRIORITY APPLIC. NO.: JP 2003434868 APPLIC. DATE: 20031226

NATIONAL APPLIC. NO.: JP 2003434868 APPLIC. DATE: 20031226

LANGUAGE: Japanese

ABSTRACT: DERWENT ABSTRACT: NOVELTY - A protease (I) having high degradation activity with respect to skin desmosome, in comparison with skin keratin, is new. DETAILED DESCRIPTION - An INDEPENDENT CLAIM is included for a skin cleaning agent composition (II) comprising (I). BIOTECHNOLOGY - Preferred Protease: (I) Exhibits a degradation active value, in a ratio of 1:2-3:6-7:8-9, with respect to synthetic substrates such as Ala-Ala-Ala, Ala-Pro-Ala, Ala-Ala-Pro-Phe, and Ala-Ala-Pro-Leu, at 30 degreesC for 1 minute. (I) Exhibits a degradation active value of 1, with respect to a synthetic substrate such as Ala-Ala-Ala at 30 degreesC for 1 minute, and a degradation active value of 6-9 with respect to casein at 30 degreesC for 1 minute. USE - (I) Is useful for preparing a skin cleaning agent composition (claimed). (I) Or (II) is useful for removing desquamation that spoils the fine region in rough skin. ADVANTAGE - (I) Does not exfoliate or peel the keratic layer of normal skin, excessively. (I) Does not cause skin irritations such as redness, itchiness and pain, and can be used daily. EXAMPLE - The microorganism *Bacillus* sp. HH192B strain was inoculated into Bouillon culture medium comprising sodium carbonate (1 %), and subjected to cultivation at 30 degreesC, overnight, with shaking. The culture supernatant was then ultrafiltered, and ammonium sulfate or acetone precipitation was performed. Then, the obtained product was subjected to diethylaminoethyl (DEAE) ion-exchange column chromatography using hydroxyapatite resin, and a crude refined enzyme with a purification degree of 70 % was obtained. To obtain the active ingredient, the obtained elute was centrifuged and a precipitate was collected. The precipitate was dissolved in buffer and column chromatography was performed using DEAE and hydroxyapatite. An active enzyme product (protease) was thus obtained. (15 pages)

DESCRIPTORS: *Bacillus* protease isol., purification, fermentation, bouillon, culture medium, ionexchange chromatography, skin desmosome degradation, appl., cosmetic comp. bacterium enzyme (24, 34)

SECTION: PHARMACEUTICALS-Other Pharmaceuticals-BIOMANUFACTURING and BIOCATALYSIS-Fermentation; BIOMANUFACTURING and BIOCATALYSIS-Biocatalyst Isolation and Characterization

Novel protease having high degradation activity with respect to skin desmosome, in comparison with skin keratin, useful for preparing a skin cleaning agent composition - isolation and purification of a protease from...

...ABSTRACT: protease (I) having high degradation activity with respect to

skin desmosome, in comparison with skin keratin, is new. DETAILED DESCRIPTION - An INDEPENDENT CLAIM is included for a skin cleaning agent composition...

...performed. Then, the obtained product was subjected to diethylaminoethyl (DEAE) ion-exchange column chromatography using hydroxyapatite resin, and a crude refined enzyme with a purification degree of 70 % was obtained. To...

... collected. The precipitate was dissolved in buffer and column chromatography was performed using DEAE and hydroxyapatite . An active enzyme product (protease) was thus obtained.(15 pages)

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$0.17 Estimated cost File41
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        $2.00  1 Type(s) in Format  9
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$14.92 Estimated cost File73
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$0.09 Estimated cost File91
    $0.22    0.062 DialUnits File94
        $4.05  3 Type(s) in Format  9
    $4.05  3 Types
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